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(57) Abstract

Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.

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GROUP B STREPTOCOCCUS ANTIGENS

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FIELD OF THE INVENTION

The present invention is related to antigens, more particularly protein antigens of group B streptococcus (GBS)

10 bacterial pathogen which are useful as vaccine components for therapy and/or prophylaxis.

BACKGROUND OF THE INVENTION

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Streptococcus are gram (+) bacteria that are differentiated by group specific carbohydrate antigens A through O found on their cell surface. Streptococcus groups are further distinguished by type-specific capsular polysaccharide antigens. Several serotypes have been identified for the Group B streptococcus (GBS): Ia, Ib, II, III, IV, V, VI, VII and VIII. GBS also contains antigenic proteins known as "C-proteins" (alpha, beta, gamma and delta), some of which have been cloned.

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Although GBS is a common component of the normal human vaginal and colonic flora this pathogen has long been recognized as a major cause of neonatal sepsis and meningitis, late-onset meningitis in infants, postpartum endometritis as well as mastitis in dairy herds. Expectant mothers exposed to GBS are at risk of postpartum infection and may transfer the infection to their baby as the child passes through the birth canal. Although the organism is sensitive to antibiotics, the high attack rate and rapid onset of sepsis in neonates and meningitis in infants results in high morbidity and mortality.

To find a vaccine that will protect individuals from GBS infection, researches have turned to the type-specific antigens. Unfortunately these polysaccharides have proven to be poorly immunogenic in humans and are restricted to the particular serotype from which the polysaccharide originates. Further, capsular polysaccharide elicit a T cell independent response i.e. no IgG production.

Consequently capsular polysaccharide antigens are unsuitable as a vaccine component for protection against GBS infection.

Others have focused on the C-protein beta antigen which demonstrated immunogenic properties in mice and rabbit models. This protein was found to be unsuitable as a human vaccine because of its undesirable property of interacting with high affinity and in a non-immunogenic manner with the Fc region of human IgA. The C-protein alpha antigen is rare in type III serotypes of GBS which is the serotype responsible for most GBS mediated conditions and is therefore of little use as a vaccine component.

Therefore there remains an unmet need for GBS antigens that may be used as vaccine components for the prophylaxis and/or therapy of GBS infection.

SUMMARY OF THE INVENTION

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According to one aspect, the present invention provides an isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide comprising a sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,

SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,

SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,

SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

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SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
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In other aspects, there is provided vectors comprising polynucleotides of the invention operably linked to an expression control region, as well as host cells transfected with said vectors and methods of producing polypeptides comprising culturing said host cells under conditions suitable for expression.

In yet another aspect, there is provided novel polypeptides encoded by polynucleotides of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

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Figure 1a is the DNA sequence of clone 1 (SEQ ID NO :1) with corresponding amino acid sequences for open reading frames; figure 1b is the amino acid sequence SEQ ID NO: 2; figure 1c is the amino acid sequence SEQ ID NO: 3; figure 1d is the amino acid sequence SEQ ID NO: 4; figure 1e is the amino acid sequence SEQ ID NO: 5; figure 1f is the amino acid sequence SEQ ID NO: 6;
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Figure 2a is the DNA sequence of clone 2 (SEQ ID NO :7) with

corresponding amino acid sequences for open reading frames;
figure 2b is the amino acid sequence SEQ ID NO: 8;
figure 2c is the amino acid sequence SEQ ID NO: 9;
figure 2d is the amino acid sequence SEQ ID NO:10;
figure 2e is the amino acid sequence SEQ ID NO:11;

figure 2f is the amino acid sequence SEQ ID NO:12;

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Figure 3a is the DNA sequence of clone 3 (SEQ ID NO :13)
    with corresponding amino acid sequences for open reading
    frames;
    figure 3b is the amino acid sequence SEQ ID NO:14;
    figure 3c is the amino acid sequence SEQ ID NO:15;
    figure 3d is the amino acid sequence SEQ ID NO:16;
    figure 3e is the amino acid sequence SEQ ID NO:17;
    figure 3f is the amino acid sequence SEQ ID NO:18;
    figure 3g is the amino acid sequence SEQ ID NO:19;
    figure 3h is the amino acid sequence SEQ ID NO:20;
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    figure 3i is the amino acid sequence SEQ ID NO:21;
    Figure 4a is the DNA sequence of clone 4 (SEQ ID NO :22)
    with corresponding amino acid sequences for open reading
15
    frames;
    figure 4b is the amino acid sequence SEQ ID NO:23;
    figure 4c is the amino acid sequence SEQ ID NO:24;
    figure 4d is the amino acid sequence SEQ ID NO:25;
    figure 4e is the amino acid sequence SEQ ID NO:26;
20
    Figure 5a is the DNA sequence of clone 5 (SEQ ID NO :27)
    with corresponding amino acid sequences for open reading
    frames;
    figure 5b is the amino acid sequence SEQ ID NO:28;
    figure 5c is the amino acid sequence SEQ ID NO:29;
    figure 5d is the amino acid sequence SEQ ID NO:30;
    figure 5e is the amino acid sequence SEQ ID NO:31;
    Figure 6a is the DNA sequence of clone 6 (SEQ ID NO :32) ;
    figure 6b is the amino acid sequence SEQ ID NO:33;
    figure 6c is the amino acid sequence SEQ ID NO:34;
    figure 6d is the amino acid sequence SEQ ID NO:35;
    figure 6e is the amino acid sequence SEQ ID NO:36;
    Figure 7a is the DNA sequence of clone 7 (SEQ ID NO :37);
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    figure 7b is the amino acid sequence SEQ ID NO:38;
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figure 7c is the amino acid sequence SEQ ID NO:39; figure 7d is the amino acid sequence SEQ ID NO:40; figure 7e is the amino acid sequence SEQ ID NO:41;
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5 Figure 8 is the DNA sequence of a part of clone 7 including a signal sequence (SEQ ID NO :42);

Figure 9 is the DNA sequence of a part of clone 7 without a signal sequence (SEQ ID NO :43);

10 Figure 9a is the amino acid sequence (SEQ ID NO:44);

Figure 10 represents the distribution of anti-GBS ELISA titers in sera from CD-1 mice immunized with recombinant GBS protein corresponding to the SEQ ID NO:39.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to novel antigenic polypeptides of group B streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,

SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,

SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,

10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,

SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,

SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,

SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,

SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,

15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

A preferred embodiment of the invention includes SEQ ID NO :39 and SEQ ID NO:44.

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A further preferred embodiment of the invention is SEQ ID NO :39.

A further preferred embodiment of the invention is SEQ ID NO :44.

As used herein, "fragments", "derivatives" or "analogs" of the polypeptides of the invention include those polypeptides in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably conserved) and which may be natural or unnatural.

The terms «fragments», «derivatives» or «analogues» of polypeptides of the present invention also include polypeptides which are modified by addition, deletion,

substitution of amino acids provided that the polypeptides retain the capacity to induce an immune response.

By the term «conserved amino acid» is meant a substitution of one or more amino acids for another in which the antigenic determinant (including its secondary structure and hydropathic nature) of a given antigen is completely or partially conserved in spite of the substitution.

10 For example, one or more amino acid residues within the sequence can be substituted by another amino acid of a similar polarity, which acts as a functional equivalent, resulting in a silent alteration. Substitutes for an amino acid within the sequence may be selected from other members of the class to which the amino acid belongs. For example, 15 the nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan and methionine. The polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine and glutamine. The positively charged (basic) 20 amino acids include arginine, lysine and histidine. The negatively charged (acidic) amino acids include aspartic acid and glutamic acid.

25 Preferably, derivatives and analogs of polypeptides of the invention will have about 70% identity with those sequences illustrated in the figures or fragments thereof. That is, 70% of the residues are the same. More preferably polypeptides will have greater than 95% homology. In another preferred embodiment, derivatives and analogs of polypeptides of the invention will have fewer than about 20 amino acid residue substitutions, modifications or deletions and more preferably less than 10. Preferred substitutions are those known in the art as conserved i.e. the substituted residues share physical or chemical properties such as hydrophobicity, size, charge or functional groups.

Furthermore, in those situations where amino acid regions are found to be polymorphic, it may be desirable to vary one or more particular amino acids to more effectively mimic the different epitopes of the different GBS strains.

Also included are polypeptides which have fused thereto other compounds which alter the polypeptides biological or pharmacological properties i.e. polyethylene glycol (PEG) to increase half-life; leader or secretory amino acid sequences for ease of purification; prepro- and pro- sequences; and (poly) saccharides.

Moreover, the polypeptides of the present invention can be modified by terminal -NH₂ acylation (eg. by acetylation, or thioglycolic acid amidation, terminal carbosy amidation, e.g. with ammonia or methylamine) to provide stability, increased hydrophobicity for linking or binding to a support or other molecule.

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Also contemplated are hetero and homo polypeptide multimers of the polypeptide fragments, analogues and derivatives. These polymeric forms include, for example, one or more polypeptides that have been cross-linked with cross-linkers such as avidin/biotin, gluteraldehyde or dimethyl-superimidate. Such polymeric forms also include polypeptides containing two or more tandem or inverted contiguous sequences, produced from multicistronic mRNAs generated by recombinant DNA technology.

Preferably, a fragment, analog or derivative of a polypeptide of the invention will comprise at least one antigenic region i.e. at least one epitope.

In order to achieve the formation of antigenic polymers

(i.e. synthetic multimers), polypeptides may be utilized having bishaloacetyl groups, nitroarylhalides, or the like,

where the reagents being specific for thio groups. Therefore, the link between two mercapto groups of the different peptides may be a single bond or may be composed of a linking group of at least two, typically at least four, and not more than 16, but usually not more than about 14 carbon atoms.

In a particular embodiment, polypeptide fragments, analogs and derivatives of the invention do not contain a methionine (Met) starting residue. Preferably, polypeptides will not incorporate a leader or secretory sequence (signal sequence). The signal portion of a polypeptide of the invention may be determined according to established molecular biological techniques. In general, the polypeptide of interest may be isolated from a GBS culture and subsequently sequenced to determine the initial residue of the mature protein and therefor the sequence of the mature polypeptide.

- According to another aspect, there is provided vaccine compositions comprising one or more GBS polypeptides of the invention in admixture with a pharmaceutically acceptable carrier diluent or adjuvant.
- Suitable adjuvants include oils i.e. Freund's complete or incomplete adjuvant; salts i.e. AlK(SO₄)₂, AlNa(SO₄)₂, AlNH₄(SO₄)₂, Al(OH)₃, AlPO₄, silica, kaolin; saponin derivative; carbon polynucleotides i.e. poly IC and poly AU and also detoxified cholera toxin (CTB) and E.coli heat labile toxin for induction of mucosal immunity. Preferred adjuvants include QuilATM, AlhydrogelTM and AdjuphosTM. Vaccines of the invention may be administered parenterally by injection, rapid infusion, nasopharyngeal absorption, dermoabsorption, or bucal or oral.

Vaccine compositions of the invention are used for the treatment or prophylaxis of streptococcus infection and/or diseases and symptoms mediated by streptococcus infection, 5 in particular group A streptococcus (pyogenes), group B streptococcus (GBS or agalactiae), dysgalactiae, uberis, nocardia as well as Staphylococcus aureus. General information about Streptococcus is available in Manual of Clinical Microbiology by P.R.Murray et al. (1995, 6th Edition, 10 ASM Press, Washington, D.C.). More particularly group B streptococcus, agalactiae. In a particular embodiment vaccines are administered to those individuals at risk of GBS infection such as pregnant women and infants for sepsis, meningitis and pneumonia as well as immunocompromised 15 individuals such as those with diabetes, liver disease or cancer. Vaccines may also have veterinary applications such as for the treatment of mastitis in cattle which is mediated by the above mentioned bacteria as well as E.coli.

The vaccine of the present invention can also be used for the manufacture of a medicament used for the treatment or prophylaxis of streptococcus infection and/or diseases and symptoms mediated by streptococcus infection, in particular group A streptococcus (pyogenes), group B streptococcus (GBS or agalactiae), dysgalactiae, uberis, nocardia as well as Staphylococcus aureus. More particularly group B streptococcus, agalactiae.

Vaccine compositions are preferably in unit dosage form of about 0.001 to 100 µg/kg (antigen/body weight) and more preferably 0.01 to 10 µg/kg and most preferably 0.1 to 1 µg/kg 1 to 3 times with an interval of about 1 to 12 weeks intervals between immunizations, and more preferably 1 to 6

weeks.

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According to another aspect, there is provided polynucleotides encoding polypeptides of group B

- 5 streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:
 - SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
 - SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
 - SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
- 10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
 - SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
 - SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
 - SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
 - SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
- 15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9 (SEQ ID NO: 43) which correspond to the open reading frames, encoding polypeptides of the invention.

- Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9(SEQ ID NO: 43) and fragments, analogues and derivatives thereof.
 - More preferred polynucleotides of the invention are those illustrated in Figures 7 (SEQ ID NO: 37), 8 (SEQ ID NO: 42) and 9 (SEQ ID NO: 43).
- Most preferred polynucleotides of the invention are those illustrated in Figures 8 (SEQ ID NO : 42) and 9(SEQ ID NO :

43).

It will be appreciated that the polynucleotide sequences illustrated in the figures may be altered with degenerate codons yet still encode the polypeptides of the invention.

Due to the degeneracy of nucleotide coding sequences, other polynucleotide sequences which encode for substantially the same polypeptides of the present invention may be used in the practice of the present invention. These include but are not limited to nucleotide sequences which are altered by the substitution of different codons that encode the same amino acid residue within the sequence, thus producing a silent change.

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Accordingly the present invention further provides polynucleotides which hybridize to the polynucleotide sequences herein above described (or the complement sequences thereof) having 50% and preferably at least 70% identity between sequences. More preferably polynucleotides are hybridizable under stringent conditions i.e. having at least 95% identity and most preferably more than 97% identity.

By capable of hybridizing under stringent conditions is meant annealing of a nucleic acid molecule to at least a region of a second nucleic acid sequence (whether as cDNA, mRNA, or genomic DNA) or to its complementary strand under standard conditions, e.g. high temperature and/or low salt content, which tend to disfavor hybridization of noncomplementary nucleotide sequences. A suitable protocol is described in Maniatis T. et al., Molecular cloning: A Laboratory Manual, Cold Springs Harbor Laboratory, 1982, which is herein incorporated by reference.

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In a further aspect, polynucleotides encoding polypeptides

of the invention, or fragments, analogs or derivatives thereof, may be used in a DNA immunization method. That is, they can be incorporated into a vector which is replicable and expressible upon injection thereby producing the antigenic polypeptide in vivo. For example polynucleotides may be incorporated into a plasmid vector under the control of the CMV promoter which is functional in eukaryotic cells. Preferably the vector is injected intramuscularly.

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According to another aspect, there is provided a process for producing polypeptides of the invention by recombinant techniques by expressing a polynucleotide encoding said polypeptide in a host cell and recovering the expressed polypeptide product. Alternatively, the polypeptides can be 15 produced according to established synthetic chemical techniques i.e. solution phase or solid phase synthesis of oligopeptides which are ligated to produce the full polypeptide (block ligation).

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For recombinant production, host cells are transfected with vectors which encode the polypeptide, and then cultured in a nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes. Suitable vectors are those that are viable and replicable in the chosen host and include chromosomal, non-chromosomal and synthetic DNA sequences e.g. bacterial plasmids, phage DNA, baculovirus, yeast plasmids, vectors derived from combinations of plasmids and phage DNA. The polypeptide sequence may be incorporated in the vector at the appropriate site using restriction enzymes such that it is operably linked to an expression control region comprising a promoter, ribosome binding site (consensus region or Shine-Dalgarno sequence), and optionally an operator (control element). One can select individual components of the 35 expression control region that are appropriate for a given

host and vector according to established molecular biology principles (Sambrook et al, Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor, N.Y., 1989 incorporated herein by reference). Suitable promoters include but are not limited to LTR or SV40 promoter, E.coli lac, tac or trp promoters and the phage lambda P. promoter. Vectors will preferably incorporate an origin of replication as well as selection markers i.e. ampicillin resistance gene. Suitable bacterial vectors include pET, pQE70, pQE60, pQE-9, pbs, pD10 phagescript, psiX174, pbluescript SK, pbsks, pNH8A, pNH16a, pNH18A, pNH46A, ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 and eukaryotic vectors pBlueBacIII, pWLNEO, pSV2CAT, pOG44, pXT1, pSG, pSVK3, pBPV, pMSG and pSVL. Host cells may be bacterial i.e. E.coli, Bacillus subtilis,

15 Streptomyces; fungal i.e. Aspergillus niger, Aspergillus nidulins; yeast i.e. Saccharomyces or eukaryotic i.e. CHO, COS.

Upon expression of the polypeptide in culture, cells are 20 typically harvested by centrifugation then disrupted by physical or chemical means (if the expressed polypeptide is not secreted into the media) and the resulting crude extract retained to isolate the polypeptide of interest. Purification of the polypeptide from culture media or lysate may be achieved by established techniques depending on the 25 properties of the polypeptide i.e. using ammonium sulfate or ethanol precipitation , acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, hydroxylapatite 30 chromatography and lectin chromatography. purification may be achieved using HPLC.

The polypeptide may be expressed with or without a leader or secretion sequence. In the former case the leader may be removed using post-translational processing (see US

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4,431,739; 4,425,437; and 4,338,397 incorporated herein by reference) or be chemically removed subsequent to purifying the expressed polypeptide.

- According to a further aspect, the GBS polypeptides of the invention may be used in a diagnostic test for streptococcus infection in particular GBS infection. Several diagnostic methods are possible, for example detecting streptococcus organism in a biological sample, the following procedure may be followed:
 - a) obtaining a biological sample from a patient;
 - b) incubating an antibody or fragment thereof reactive with a GBS polypeptide of the invention with the biological sample to form a mixture; and
- 15 c) detecting specifically bound antibody or bound fragment in the mixture which indicates the presence of streptococcus.

Alternatively, a method for the detection of antibody
20 specific to a streptococcus antigen in a biological sample
containing or suspected of containing said antibody may be
performed as follows:

a) isolating a biological sample from a patient;

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- b) incubating one or more GBS polypeptides of the 25 invention or fragments thereof with the biological sample to form a mixture; and
 - c) detecting specifically bound antigen or bound fragment in the mixture which indicates the presence of antibody specific to streptococcus.

One of skill in the art will recognize that this diagnostic test may take several forms, including an immunological test such as an enzyme-linked immunosorbent assay (ELISA), a radioimmunoassay or a latex agglutination assay, essentially to determine whether antibodies specific for the protein are present in an organism.

The DNA sequences encoding polypeptides of the invention may also be used to design DNA probes for use in detecting the presence of streptococcus in a biological sample suspected of containing such bacteria. The detection method of this invention comprises:

- a) isolating the biological sample from a patient;
- b) incubating one or more DNA probes having a DNA sequence encoding a polypeptide of the invention or fragments thereof with the biological sample to form a mixture; and
- c) detecting specifically bound DNA probe in the mixture which indicates the presence of streptococcus bacteria.
- The DNA probes of this invention may also be used for detecting circulating streptococcus i.e. GBS nucleic acids in a sample, for example using a polymerase chain reaction, as a method of diagnosing streptococcus infections. The probe may be synthesized using conventional techniques and may be immobilized on a solid phase, or may be labeled with a detectable label. A preferred DNA probe for this application is an oligomer having a sequence complementary to at least about 6 contiguous nucleotides of the GBS polypeptides of the invention.

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Another diagnostic method for the detection of streptococcus in a patient comprises:

- a) labeling an antibody reactive with a polypeptide of the invention or fragment thereof with a detectable label;
- 30 b) administering the labeled antibody or labeled fragment to the patient; and
 - c) detecting specifically bound labeled antibody or labeled fragment in the patient which indicates the presence of streptococcus.

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A further aspect of the invention is the use of the GBS

polypeptides of the invention as immunogens for the production of specific antibodies for the diagnosis and in particular the treatment of streptococcus infection. Suitable antibodies may be determined using appropriate screening methods, for example by measuring the ability of a particular antibody to passively protect against streptococcus infection in a test model. One example of an animal model is the mouse model described in the examples herein. The antibody may be a whole antibody or an antigen-10 binding fragment thereof and may in general belong to any immunoglobulin class. The antibody or fragment may be of animal origin, specifically of mammalian origin and more specifically of murine, rat or human origin. It may be a natural antibody or a fragment thereof, or if desired, a 15 recombinant antibody or antibody fragment. The term recombinant antibody or antibody fragment means antibody or antibody fragment which were produced using molecular biology techniques. The antibody or antibody fragments may be polyclonal, or preferably monoclonal. It may be specific for a number of epitopes associated with the GBS 20 polypeptides but is preferably specific for one.

EXAMPLE 1 Murine model of lethal Group B Streptococcus (GBS)
25 infection

The mouse model of GBS infection is described in detail in Lancefield et al (J Exp Med 142:165-179,1975). GBS strain C388/90 (Clinical isolate obtained in 1990 from the cephalorachidian fluid of a patient suffering from meningitis, Children's Hospital of Eastern Ontario, Ottawa, Canada) and NCS246 (National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Edmonton, Canada) were respectively serotyped as type Ia/c and type II/R.

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To increase their virulence, the GBS strains C388/90 (serotype Ia/c) and NCS 246 (serotype II/R) were serially passaged through mice as described previously (Lancefield et al. J Exp Med 142:165-179, 1975). Briefly, the increase of virulence was monitored using intraperitoneal inoculations of serial dilutions of a subculture in Todd-Hewitt broth obtained from either the blood or spleen of infected mice. After the last passage, infected blood samples were used to inoculate Todd-Hewitt broth. After an incubation of 2 hours at 37°C with 7% CO2, glycerol at a final concentration of 10 10% (v/v) was added to the culture. The culture was then aliquoted and stored at -80° C for use in GBS challenge experiments. The number of cfu of GBS present in these frozen samples was determined. The bacterial concentration necessary to kill 100% (LD100) of the 18 weeks old mice were 15 determined to be 3.5X10⁵ and 1.1X10⁵ respectively for GBS strain C388/90 and NCS246, which corresponded to a significant increase in virulence for both strains. Indeed, the LD100 recorded before the passages for these two strains was higher than 10° cfu. 20

In a bacterial challenge, a freshly thawed aliquot of a virulent GBS strain was adjusted to the appropriate bacterial concentration using Todd-Hewitt broth and 1ml was injected intraperitoneally to each female CD-1 mouse. 25 mice used for the passive protection experiments were 6 to 8 weeks old, while the ones used for the active protection experiments were approximately 18 weeks old at the time of the challenge. All inocula were verified by colony counts. 30 Animals were observed for any sign of infection four times daily for the first 48 h after challenge and then daily for the next 12 days. At the end of that period, blood samples were obtained from the survivors and frozen at -20°C. spleen obtained from each mouse that survived the challenge was cultured in order to identify any remaining GBS. 35

EXAMPLE 2 Immunization and protection in mice with formaldehyde killed whole GBS cells

Formaldehyde killed GBS whole cells were prepared according to the procedures described in Lancefield et al (J Exp Med 142:165-179,1975). Briefly, an overnight culture on sheep blood agar plates (Quelab Laboratories, Montreal, Canada) of a GBS strain was washed twice in PBS buffer (phosphate buffered-saline, pH7.2), adjusted to approximately 3X10° cfu/mL and incubated overnight in PBS containing 0.3% (v/v) formaldehyde. The killed GBS suspension was washed with PBS

and kept frozen at -80°C.

- 15 Female CD-1 mice, 6 to 8 weeks old (Charles River, St-Constant, Québec, Canada), were injected subcutaneously three times at two weeks interval with 0.1 ml of formaldehyde killed cells of GBS strain C388/90 (~6X107 GBS), or 0.1 ml of PBS for the control group. On the day before the immunization, AlhydrogelTM (Superfos Biosector, Frederikssund, Denmark) at a final concentration of 0.14 mg or 0.21 mg of Al, was added to these preparations and incubated overnight at 4°C with agitation. Serum samples were obtained from each mouse before the beginning of the immunization protocol and two weeks after the last injection. The sera were frozen at -20°C.
- Eight mice in each control group injected with PBS and the group immunized with formaldehyde killed whole cells GBS strain C388/90 (Ia/c) were challenged with 1.5X10⁴ cfu of GBS strain C388/90 (Ia/c) one week after the third injection. All mice immunized with the formaldehyde killed GBS whole cells survived the homologous challenge while, within 5 days after the challenge, only 4 out of the 8 mice injected with PBS survived from the infection. In order to increase the mortality rate in the control groups, the

bacterial suspension had to be adjusted according to the age of the mice at the time of the bacterial challenge. In subsequent challenge experiments, when mice were older than 15 weeks, the bacterial inoculum was increased to concentrations between 3.0X10⁵ and 2.5X10⁶ cfu.

Table 1 Immunization of CD1 mice with formaldehyde killed whole cells of GBS and subsequent homologous challenge [strain C388/90 and heterologous challenge [strain NCS246 (II/R)].

antigenic preparations used for immunization ¹	number of living mice 14 days after the bacterial challenge (% Survival)				
	homologous challenge: strain C388/90 (Ia/c)	heterologous challenge: strain NCS246 (II/R)			
1st infection					
formaldehyde killed cells of GBS strain C388/90 (la/c) ²	8/8 (100) ³	n.d. ⁵			
control PBS	4/8 (50)	n.d.			
2nd infection					
formaldehyde killed cells of GBS strain C388/90 (la/c)	6/6 (100)⁴	0/6 (0) ⁶			
control PBS	2/6 (33)	0/6 (0)			

¹ alhydrogel™ at a final concentration of 0.14 mg or 0.21mg of Al was used;

² approximately 6X10⁷ cfu;

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In another experiment, one group of 12 mice corresponding to a control group was injected with PBS, while a second group of 12 mice was immunized with formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Six mice from each of these two groups were challenged with 2.1X106 cfu of the GBS strain C388/90 (Ia/c) (Table I). As the first challenge experiment, all mice immunized with the GBS strain C388/90 (Ia/c) survived the homologous challenge. Only two out of the 6 mice injected with PBS survived the infection.

³ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (la/c) suspension adjusted to 1.5X10⁴ cfu;

⁴ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (la/c) suspension adjusted to 2.1X10⁶ cfu; ⁵ not done;

⁶ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS NCS246 (II/R) suspension adjusted to 1.2X10⁵ cfu.

The remaining 6 mice in both groups were then used one week later to verify whether this antigenic preparation could confer cross protection against strain NCS246 (II/R) which produce a serologically distinct capsule. None of the mice infected with this second GBS strain survived the infection. The later result suggested that most of the protective immune response induced by formaldehyde killed strain C388/90 is directed against the capsular polysaccharide and that it could be restricted to strains of that particular serotype. These results clearly indicated that this particular model of infection can be efficiently used to study the protection conferred by vaccination.

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EXAMPLE 3 Immunization of rabbit with formaldehyde killed whole GBS cells and passive protection in mice

A New Zealand rabbit (2.5 kg, Charles River, St-Constant, Québec, Canada) was immunized with formaldehyde killed cells of GBS strain C388/90 (Ia/c) to obtain hyperimmune serum. This rabbit was injected subcutaneously three times at three weeks interval with approximately 1.5X10° cfu of formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Freund's complete adjuvant (Gibco BRL Life Technologies, Grand Island, New York) was used as the adjuvant for the first immunization, while Freund's incomplete adjuvant (Gibco BRL) was used for the following two injections. Serum samples were obtained before the beginning of the immunization protocol and two weeks after the last injection. The sera were frozen at -20°C.

The ability of this particular rabbit hyperimmune serum to passively protect mice against a lethal infection with GBS

was also evaluated. Intraperitoneal injection of mice with either 15 or 25 μ L of hyperimmune rabbit serum 18 hours before the challenge protected 4 out of 5 mice (80%) against the infection. Comparatively, survival rates lower than 20% were recorded for mice in the control group injected with PBS or serum obtained from a rabbit immunized with meningococcal outer membrane preparation. This result clearly indicates that the immunization of another animal species with killed GBS cells can induce the production of antibodies that can passively protect mice. This reagent will also be used to characterize clones.

Table 2 Passive protection of CD-1 mice conferred by rabbit serum obtained after immunization with formaldehyde killed group B whole streptococci (strain C388/90 (Ia/c)) antigenic preparation

groups	number of living mice 14 days after the bacterial challenge with GBS strain C388/90 (Ia/c) ²	% survival
rabbit hyperimmune serum² - 25 μl	4/5	80
rabbit hyperimmune serum¹ - 15 μl	4/5	80
control rabbit serum - 25 μl	1/5	20
control PBS	1/10	10

Freund's complete adjuvant was used for first immunization, and Freund's incomplete adjuvant for the following two injections;

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intraperitoneal challenge with 1 ml Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2X10⁴ cfu.

EXAMPLE 4 Recombinant production of His.Tag-GBS fusion protein

The coding region of a GBS gene was amplified by PCR (DNA Thermal Cycler GeneAmp PCR system 2400 Perkin Elmer, San Jose, CA) from the genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BqlII (AGATCT) and HindIII (AAGCTT), respectively. The PCR product was purified from 10 agarose gel using a Qiaex II gel extraction kit from Oiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII (Pharmacia Canada Inc Baie d'Urfe, Canada), and extracted with phenol:chloroform before ethanol precipitation. The pET-32b(+) vector (Novagen, Madison, WI) 15 containing the thioredoxin-His. Tag sequence was digested with the restriction enzymes BqlII and HindIII, extracted with phenol:chloroform, and then ethanol precipitated. BglII-HindIII genomic DNA fragment was ligated to the BglII-HindIII pET-32b(+) vector to create the coding sequence for 20 thioredoxin-His. Tag-GBS fusion protein whose gene was under control of the T7 promoter. The ligated products were transformed into E. coli strain XLI Blue MRF' $(\Delta (mcrA) 183\Delta$ (mcrCB-hsdSMR-mrr)173 endAl supE44 thi-1 recAl qyrA96 relAl lac [F'proAB lacIqZΔM15Tn10 (Tetr)]c) (Stratagene, La Jolla, 25 CA) according to the method of Simanis (Hanahan, D. DNA Cloning, 1985, D.M. Glover (ed.), pp. 109-135). recombinant pET plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing (Taq Dye Deoxy 30 Terminator Cycle Sequencing kit, ABI, Foster City, CA). recombinant pET plasmid was transformed by electroporation (Gene Pulser II apparatus, BIO-RAD Labs, Mississauga, Canada) into E. coli strain AD494 (DE3) (Δara leu7697 ΔlacX74 ΔphoA PvuII phoR ΔmalF3 F'[lac⁺(lacIq) pro]

trxB::Kan (DE3)) (Novagen, Madison, WI). In this strain of

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E. coli, the T7 promoter controlling expression of the fusion protein, is specifically recognized by the T7 RNA polymerase (present on the $\lambda DE3$ prophage) whose gene is under the control of the lac promoter which is inducible by isopropyl- β -D-thio-galactopyranoside (IPTG).

The transformant AD494 (DE3)/rpET was grown at 37°C with agitation at 250 rpm in LB broth (peptone 10g/L, Yeast extract 5g/L, NaCl 10g/L) containing 100 μ g of ampicillin (Sigma-Aldrich Canada Ltd., Oakville, Canada) per mL until the A₆₀₀ reached a value of 0.6. In order to induce the production of the thioredoxin-His.Tag-GBS fusion protein, the cells were incubated for 2 additional hours in the presence of IPTG at a final concentration of 1mM. The bacterial cells were harvested by centrifugation.

The recombinant fusion protein produced by AD494(DE3)/rpET32 upon IPTG induction for 2h was partially obtained as insoluble inclusion bodies which were purified from 20 endogenous E. coli proteins by the isolation of insoluble aggregates (Gerlach, G.F. et al 1992, Infect. Immun. 60:892). Induced cells from a 500 mL culture were resuspended in 20 mL of 25% sucrose-50mM Tris-HCl buffer (pH8.0) and frozen at -70°C. Lysis of cells in thawed 25 suspension was achieved by the addition of 5mL of a solution of lysozyme (10mg/mL) in 250mM Tris-HCl buffer (pH8.0) followed by an incubation of 10 to 15 min on ice, and the addition of 150mL of detergent mix (5 parts of 20mM Tris-HCl buffer [pH7.4]-300mM NaCl-2% deoxycholic acid-2% Nonidet P-30 40 and 4 parts of 100mM Tris-HCl buffer [pH8]-50mM EDTA-2% Triton X-100) followed by 5 min incubation on ice. sonication, protein aggregates were harvested by centrifugation for 30 min at 35,000 X g and a sample of the soluble cellular fraction was kept. The aggregated proteins 35 were solubilized in 6M guanidine hydrochloride. The

presence of the fusion protein in both the soluble and insoluble fractions was shown by Western Blot analysis using the serum of a mouse injected with formaldehyde killed cells of GBS strain C388/90 (Ia/c) that survived a bacterial challenge with the corresponding GBS strain.

The purification of the fusion protein from the soluble fraction of IPTG-induced AD494 (DE3) / rpET was done by affinity chromatography based on the properties of the His. Tag sequence (6 consecutive histidine residues) to bind 10 to divalent cations (Ni²⁺) immobilized on the His.Bind metal chelation resin (Novagen, Madison, WI). The purification method used are those described in the pET system Manual, 6th Edition (Novagen, Madison, WI). Briefly, the pelleted cells obtained from a 100mL culture induced with IPTG was resuspended in 4mL of Binding buffer (5mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9), sonicated, and spun at 39,000 X g for 20 min to remove debris. The supernatant was filtered $(0.45\mu\text{m}\text{ pore size membrane})$ and deposited on a column of 20 His.Bind resin equilibrated in Binding buffer. The column was then washed with 10 column volumes of Binding buffer followed by 6 column volumes of Wash buffer (20mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The thioredoxin-His.Tag-GBS fusion protein was eluted with Elute buffer (1M 25 imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The removal of the salt and imidazole from the sample was done by dialysis against 3 X 1 liter PBS at 4°C.

The quantities of fusion protein obtained from either the soluble or insoluble cytoplasmic fractions of *E. coli* were estimated by Coomassie staining of a sodium dodecyl sulfate (SDS)-polyacrylamide gel with serial dilutions of these proteins and a bovine serum albumin standard (Pierce Chemical Co. Rockford, Ill.).

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EXAMPLE 5 Recombinant production of GBS protein under control of lambda P_L promoter

The DNA coding region of a GBS protein was inserted downstream of the promoter $\lambda P_{\rm L}$ into the translation vector pURV22. This plasmid was derived from p629 (George et al, 1987, Bio/Technology 5:600) from which the coding region for a portion of the herpes simplex virus type I (HSV-I) glycoprotein (gD-1) was removed and the ampicillin 10 resistance gene replaced by a kanamycin cassette obtained from the plasmid vector pUC4K (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada). The vector contained a cassette of the bacteriophage λ cI857 temperature sensitive repressor gene from which the functional $P_{\text{\tiny R}}$ promoter had been deleted. 15 The inactivation of the cI857 repressor by temperature increase from the ranges of 30-37°C to 37-42°C resulted in the induction of the gene under the control of λ P_L. The translation of the gene was controlled by the ribosome binding site cro followed downstream by a BglII restriction 20 site (AGATCT) and the ATG: ACTAAGGAGGTTAGATCTATG.

Restriction enzymes and T4 DNA ligase were used according to suppliers (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada; and New England Biolabs Ltd., Mississauga, Canada). 25 Agarose gel electrophoresis of DNA fragments was performed as described by Sambrook et al. (Molecular cloning : A laboratory Manual, 1989, Cold Spring Harbor Laboratory Press, N.Y). Chromosomal DNA of the GBS bacteria was prepared according to procedures described in Jayarao et al (J. Clin. Microbiol., 1991, 29:2774). DNA amplification 30 reactions by polymerase chain reaction (PCR) were made using DNA Thermal Cycler GeneAmp PCR system 2400 (Perkin Elmer, San Jose, CA). Plasmids used for DNA sequencing were purified using plasmid kits from Qiagen (Chatsworth, CA). DNA fragments were purified from agarose gels using Qiaex II 35

gel extraction kits from Qiagen (Chatsworth, CA). Plasmid transformations were carried out by the method described by Hanahan (DNA Cloning, Glover (ed.) pp, 109-135, 1985). The sequencing of genomic DNA inserts in plasmids was done using synthetic oligonucleotides which were synthesized by oligonucleotide synthesizer model 394 (the Perkin-Elmer Corp., Applied Biosystems Div. (ABI), Foster City, CA). The sequencing reactions were carried out by PCR using the Tag Dye Deoxy Terminator Cycle Sequencing kit (ABI, Foster City, 10 CA) and DNA electrophoresis was performed on automated DNA sequencer 373A (ABI, Foster City, CA). The assembly of the DNA sequence was performed using the program Sequencer 3.0 (Gene Codes Corporation, Ann Arbor, MI). Analysis of the DNA sequences and their predicted polypeptides was performed with the program Gene Works version 2.45 (Intelligenetics, Inc., Mountain View CA).

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The coding region of the GBS gene was amplified by PCR from GBS strain C388/90 (Ia/c) genomic DNA using oligos that 20 contained base extensions for the addition of restriction sites BglII (AGATCT) and XbaI(TCTAGA), respectively. The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and XbaI, and extracted with phenol:chloroform before ethanol precipitation. The pURV22 25 vector was digested with the restriction enzymes BglII and XbaI, extracted with phenol:chloroform, and ethanol precipitated. The BglII-XbaI genomic DNA fragment was ligated to the BglII-XbaI pURV22 vector in which the GBS gene was under the control of the λPL promoter. The ligated 30 products were transformed into E. coli strain XLI Blue MRF' $(\Delta (mcrA) 183\Delta (mcrCB-hsdSMR-mrr) 173 endA1 supE44 thi-1 recA1$ gyrA96 relA1 lac[F' proAB lac1qZAM15 Tn10(Tetr)]c) (Stratagene, La Jolla CA) according to the methods described in Hanahan, supra. Transformants harboring plasmids with the 35

insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook et al, <u>supra</u>). The recombinant pURV22 plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

The transformant XLI Blue MRF'/rpURV22 was grown at 34°C with agitation at 250 rpm in LB broth containing $50\mu g$ of kanamycin per mL until the A_{600} reached a value of 0.6. In order to induce the production of the fusion protein, the cells were incubated for 4 additional hours at 39°C. The bacterial cells were harvested by centrifugation , resuspended in sample buffer, boiled for 10 min and kept at -20°C.

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EXAMPLE 6 Subcloning GBS protein gene in CMV plasmid pCMV-GH

The DNA coding region of a GBS protein was inserted in phase downstream of the human growth hormone (hGH) gene which was under the transcriptional control of the cytomegalovirus (CMV) promoter in the plasmid vector pCMV-GH (Tang et al, Nature, 1992, 356:152). The CMV promoter is non functional in E. coli cells but active upon administration of the plasmid in eukaryotic cells. The vector also incorporated the ampicillin resistance gene.

The coding region of the gene was amplified by PCR from genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT). The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII, and extracted with phenol:chloroform before ethanol precipitation. The pCMV-GH vector (Laboratory of Dr. Stephen

A. Johnston, Department of Biochemistry, The University of Texas, Dallas, Texas) containing the human growth hormone to create fusion proteins was digested with the restriction enzymes BamHI and HindIII, extracted with phenol:chloroform, and ethanol precipitated. The 1.3-kb BglII-HindIII genomic DNA fragment was ligated to the BamHI -HindIII pCMV-GH vector to create the hGH-GBS fusion protein under the control of the CMV promoter. The ligated products were transformed into E. coli strain DH5 α [ϕ 80 lacZ Δ M15 endA1 10 recA1 hsdR17 ($^{r}K^{-m}K^{+}$) supE44 $thi-1\lambda^{-}$ gyrA96 relA1 $\Delta(lacZYA$ argF)U169] (Gibco BRL, Gaithersburg, MD) according to the methods described by Hanahan, supra. Transformants harboring plasmids with the insert were identified by analysis of lysed cells submitted to electrophoresis on 15 agarose gel (Sambrook, J. et al , supra). The recombinant pCMV plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

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EXAMPLE 7 Immunological activity of GBS protein to GBS challenge

Four groups of 12 female CD-1 mice (Charles River, St
Constant, Quebec, Canada) of 6 to 8 weeks were injected subcutaneously three times at three week intervals with 0.1mL of the following antigenic preparations: formaldehyde killed cells of GBS strain C388/90 (~6X10⁷ cfu), 20µg of thioredoxin-His.Tag-GBS fusion protein obtained from the insoluble (inclusion bodies) or 20µg of the fusion protein, affinity purified (nickel column), from the soluble cytoplasmic fraction in E.coli, or 20µg of affinity purified (nickel column) thioredoxin-His.Tag control polypeptide.

20µg of QuilATM (Cedarlane Laboratories Ltd, Hornby, Canada)

was added to each antigenic preparation as the adjuvant. Serum samples were obtained from each mouse before immunization (PB) and on days 20 (TB1), 41 (TB2) and 54 (TB3) during the immunization protocols. Sera were frozen at -20°C.

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An increase of the ELISA titers was recorded after each injection of the fusion protein indicating a good primary response and a boost of the specific humoral immune response after each of the second and third administration. At the 10 end of the immunization period, the means of reciprocal ELISA titers was 456,145 for the group immunized with $20\mu g$ of fusion protein obtained from inclusion bodies compared to 290,133 for the group of mice immunized with the protein 15 from soluble fraction in E.coli. The latter result suggests that the protein obtained from inclusion bodies could be more immunogenic than the soluble protein. Analysis of mice sera in ELISA using the affinity purified thioredoxin-His. Tag to coat plates showed that negligible antibody 20 titers are made against the thioredoxin-His. Tag portion of the fusion protein. The reactivity of the sera from mice injected with the recombinant fusion protein was also tested by ELISA against formaldehyde killed whole cells of GBS strain C388/90. The antibodies induced by immunization with 25 recombinant fusion protein also recognized their specific epitopes on GBS cells indicating that their conformation is close enough to the native streptococcal protein to induce cross-reactive antibodies.

To verify whether the immune response induced by immunization could protect against GBS infection, mice were challenged with 3.5X10⁵ cfu of GBS strains C338/90(Ia/c) and 1.2X10⁵ cfu of strain NCS246(II/R) the results of which are illustrated in tables 3 and 4 respectively. Mice immunized with control thioredoxin-His.Tag peptide were not protected against challenge with either GBS strain while those

immunized with formaldehyde killed C388/90 whole cells only provided protection against homologous challenge. The thioredoxin-His.Tag-GBS fusion protein of the invention protected mice from challenge with both GBS strains. Blood and spleen culture of these mice did not reveal the presence of any GBS.

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Table 3 Survival from GBS strain C388/90 (Ia/c) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag²	1 / 6	17
formaldehyde killed C388/90 cells³	6 / 6	100
thioredoxin-His.Tag-GBS fusion (inclusion body preparation)4	6 / 6	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ⁴	6 / 6	100

intraperitoneal administration with 1 ml Todd-Hewitt culture medium adjusted to 3.5X10⁵ cfu;

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 ²⁰μg administered; posterior legs paralyzed in surviving mouse; GBS detected in blood and spleen;
 6X10⁷ cfu administered;

 $^{^4}$ 20 μ g administered.

Table 4 Survival from GBS strain NCS246 (II/R) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag²	0 / 6	0
formaldehyde killed C388/90 cells ³	2 / 6	34
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ²	5 / 5⁴	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ²	6 / 6	100

intraperitoneal administration with 1 ml Todd-Hewitt culture medium containing GBS NCS246(II/R) suspension adjusted to 1.2X10⁵ cfu.

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EXAMPLE 8 Immunization with recombinant GBS protein confers protection against experimental GBS infection

This example illustrates the protection of mice against fatal GBS infection by immunization with the recombinant protein corresponding to the SEQ ID NO:39.

20 Groups of 10 female CD-1 mice (Charles River) were immunized subcutaneously three times at three-week intervals with 20 μg of recombinant protein purified from <u>E. coli</u> strain BLR (Novagen) harboring the recombinant pURV22 plasmid vector containing the GBS gene corresponding to SEQ ID NO:42 in presence of 20 μg of QuilATM adjuvant (Cedarlane

Laboratories Ltd, Hornby, Canada) or, as control, with

² 20µg administered;

³ 6X10⁷ cfu administered;

^{10 4} one mouse died during immunization.

QuilATM adjuvant alone in PBS. Blood samples were collected from the orbital sinus on day 1, 22 and 43 prior to each immunization and fourteen days (day 57) following the third injection. One week later the mice were challenged with approximately 10⁴ to 10⁶ CFU of various virulent GBS strains. Samples of the GBS challenge inoculum were plated on TSA/5% sheep blood agar plates to determine the CFU and to verify the challenge dose. Deaths were recorded for a period of 14 days and on day 14 post-challenge, the surviving mice were sacrificed and blood and spleen were tested for the presence of GBS organisms. The survival data are shown in table 5.

Prechallenge sera were analyzed for the presence of antibodies reactive with GBS by standard immunoassays. Elisa and immunoblot analyses indicated that immunization with recombinant GBS protein produced in *E. coli* elicited antibodies reactive with both, recombinant and native GBS protein. Antibody responses to GBS are described in Example 9.

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Table 5. Ability of recombinant GBS protein corresponding to SEQ ID NO: 39 to elicit protection against 8 diverse GBS challenge strains

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COH1-F

none

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Challenge strain Immunogen Designation No. alive: Type No. dead 1 rGBS protein C388/90 Ia/c 8:2 (P<0.0001)none 0:10 NCS 246 rGBS protein II/R 10:0 (P=0.0012)none 3:7 rGBS protein ATCC12401 Ιb 10:0 (P=0.001)none 3:7 rGBS protein NCS 535 V 10:0 (P=0.01)none 5:5 10 : 0 rGBS protein NCS 9842 VI (P<0.0001) none 0:10 rGBS protein NCS 915 III 7:3 $(P=0.0007)^2$ NCS 915-F³ 1:9 4:6 none NCS 954 rGBS protein III/R 7 **:** 3 (P=0.002)NCS 954-F 4:6 none 1:9 rGBS protein COH1 III 4:6 (P=0.0004)

3:7

0:10

All hemocultures from surviving mice were negative at day 14 20 post-challenge. Spleen cultures from surviving mice were negative except for few mice from experiment MB-11.

Groups of 10 mice per group were used, the number of mice surviving to infection and the number of dead mice are indicated. The survival curves corresponding to recombinant GBS protein-immunized animals were compared to the survival curves corresponding to mock-immunized animals using the log-rank test for nonparametric analysis.

² Comparison analysis to NCS915-F-immunized animals.

 $^{^{3}}$ Animals were immunized with formaldehyde-killed GBS in presence of QuilA $^{\text{TM}}$ adjuvant.

EXAMPLE 9 Vaccination with the recombinant GBS protein elicits an immune response to GBS

Groups of 10 female CD-1 mice were immunized subcutaneously with recombinant GBS protein corresponding to SEQ ID NO:39 as described in Example 8. In order to assess the antibody response to native GBS protein, sera from blood samples collected prior each immunization and fourteen days after the third immunization were tested for antibody reactive 10 with GBS cells by ELISA using plates coated with formaldehyde-killed GBS cells from type III strain NCS 954, type Ib strain ATCC12401, type V strain NCS 535 or type VI strain NCS 9842. The specificity of the raised antibodies for GBS protein was confirmed by Western blot analyses to 15 GBS cell extracts and purified recombinant antigens. The results shown in Figure 10 clearly demonstrate that animals respond strongly to recombinant GBS protein used as immunogens with median reciprocal antibody titers varying between 12000 and 128000, for sera collected after the third immunization, depending of the coating antigen. All 20 preimmune sera were negative when tested at a dilution of 1 :100. GBS-reactive antibodies were detectable in the sera of each animal after a single injection of recombinant GBS protein.

Example 10 Antigenic conservation of the GBS protein of the present invention

Monoclonal antibodies (MAbs) specific to the GBS protein of the present invention were used to demonstrate that this surface antigen is produced by all GBS and that it is also antigenically highly conserved.

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A collection of 68 GBS isolates was used to evaluate the 10 reactivity of the GBS-specific MAbs. These strains were obtained from the National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Canada; Centre Hospitalier Universitaire de Quebec, Pavillon CHUL, Quebec, Canada; American Type Culture Collection, USA; 15 Laboratoire de Sante Publique du Quebec, Canada; and Dept. of Infectious Disease, Children's Hospital and Medical Center, Seattle, USA. All eight Mabs were tested against the following panel of strains: 6 isolates of serotype Ia or Ia/c, 3 isolates of serotype Ib, 4 isolates of serotype II, 20 14 isolates of serotype III, 2 isolates of serotype IV, isolates of serotype V, 2 isolates of serotype VI, 2 isolates of serotype VII, 1 isolate of serotype VIII, 10 isolates that were not serotyped and 3 bovine S. agalactiae strains. MAb 3A2 was also reacted with additional GBS: 9 25 isolates of serotype Ia/c and 10 isolates of serotype V. The strains were grown overnight on blood agar plates at 37°C in an atmosphere of 5% CO_2 . Cultures were stored at - 70°C in heart infusion broth with 20% (v/v) glycerol.

30 To obtain the GBS protein-specific MAbs, mice were immunized three times at three-week intervals with 20 μg of purified recombinant GBS protein (SEQ ID NO :44) in the presence of 20% QuilATM adjuvant. Hybridoma cell lines were generated by fusion of spleen cells recovered from immunized mice with the nonsecreting SP2/O myeloma cell line as described

previously (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Hybrid clone supernatants were tested for specific antibody production by ELISA using formaldehyde inactivated GBS and purified recombinant GBS protein (SEQ ID NO :39 or 44) as coating antigen, as previously described (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Specific hybrid were cloned by limiting dilutions, expanded, and frozen in liquid nitrogen. Production of recombinant GBS protein was presented in Examples 4 & 5. Purified 10 recombinant GBS protein or formaldehyde inactivated GBS were resolved by electrophoresis by using the discontinuous buffer system of Laemmli as recommended by the manufacturer and then transfer onto nitrocellulose membrane for Western immunoblotting as described previously (Martin et al. 1992. 15 Infect. Immun. 60:2718-2725).

Western immunoblotting experiments clearly indicated that all eight MAbs recognized a protein band that corresponded to the purified recombinant GBS protein (SEQ ID NO :39).

20 These MAbs also reacted with a protein band present in every GBS isolates tested so far. The reactivity of these GBS-specific MAbs are presented in Table 6. Each MAb reacted well with all 46 GBS. In addition, these MAbs also recognized the 3 S. agalactiae strains of bovine origin that were tested. MAb 3A2 also recognized nineteen GBS; 9 isolates of serotype Ia/c and 10 of serotype V. The other MAbs were not tested against these additional strains.

These results demonstrated that the GBS protein (SEQ ID NO:39) was produced by all the 65 GBS and the three 3 S. agalactiae strains of bovine origin that were tested so far. More importantly, these results clearly demonstrated that the epitopes recognized by these eight GBS-specific MAbs were widely distributed and conserved among GBS. These results also indicated that these epitopes were not

restricted to serologically related isolates since representatives of all known GBS serotypes including the major disease causing groups were tested.

In conclusion, the data presented in this example clearly demonstrated that the GBS protein of the present invention is produced by all GBS and that it is antigenically highly conserved.

10

Reactivity of eight GBS protein-specific MAbs with different S. agalactiae strains as evaluated by Western immunoblots. Table 6.

Mabs	Num	Number of	each	serotype	Ype of	ω •	agalactiae	ciae st	strains	recognized	by the	MADs.
											F 6 6	
	Taor	Th	II	III	ΛI	>	IΛ	VII	VIII	NT(10)2	TOTAL	PONTING
	10 77	(~	(4)	(4)	(2)	(2)	(2)	(5)	(1)		(26)	(3)
	10/C (0)	(2));	,		C	C	c		C	46	m
2 A 2 1	9	m	4	4	N	7	7	7	7	7	2.	
Chi		,	_	7	2	2	2	2	٦	10	46	m
5ATZ	٥	n	#	۴	3	1				1	7 7	٠
177	2	۲	4	4	2	7	7	?	-1	⊃ T	40	7
TIPO	0		4			·	0	C	-	10	91	~
000	y	~	4	4	~	7	7	7	⊣	OT	٥ ۲	,
000		,	•		0	(c	c	-	10	46	<u>~</u>
RE11	9	m	4	4	7	7	7	7	7	7.		, ,
100		C	_	V	c	2	0	7	-	10	46	٠,
12812	٥	ŋ	ť	н	1	7	2		,	0,5	71	۲
1001	9	۳	4	4	N	~	~	7		ΩT	٠ ۲	
TOLTT		,			(,	(c	-	10	46	((1)
20G2	9	m	4	4	7	7	7	7	-4	7	2	

1 Nine additional strains of serotype Ia/c and 10 strains of serotype V were recognized by MAb 3A2. 2 These strains were not serotyped

10

WE CLAIM:

1. An isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
```

- 2. A polynucleotide according to claim 1, wherein said polynucleotide encodes a polypeptide having at least 95% identity to the second polypeptide.
- 3. An isolated polynucleotide encoding a polypeptide capable of generating antibodies having binding specificity for a polypeptide having a sequence selected from the group consisting of:

 SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

4. An isolated polynucleotide that is complementary to the polynucleotide of claim 1.

- 5. An isolated polynucleotide that is complementary to the polynucleotide of claim 3.
- 6. The polynucleotide of claim 1, wherein said polynucleotide is DNA.
- 7. The polynucleotide of claim 3, wherein said polynucleotide is DNA.
- 8. The polynucleotide of claim 1, wherein said polynucleotide is RNA.
- 9. The polynucleotide of claim 3, wherein said polynucleotide is RNA.
- 10. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of:

 SEQ ID NO: 1, SEQ ID NO: 7, SEQ ID NO: 13, SEQ ID NO: 22, SEQ ID NO: 27, SEQ ID NO: 32, SEQ ID NO: 37, SEQ ID NO: 42 and SEQ ID NO: 43 or fragments, analogues or derivatives thereof.
- 11. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of :

 SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43.
- 12. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 37.

13. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 42.

- 14. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 43.
- 15. A polynucleotide according to claim 10 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
- 16. A polynucleotide according to claim 11 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
- 17. A vector comprising the polynucleotide of claim 1, wherein said polynucleotide is operably linked to an expression control region.
- 18. A vector comprising the polynucleotide of claim 3, wherein said polynucleotide is operably linked to an expression control region.
- 19. A host cell transfected with the vector of claim 17.
- 20. A host cell transfected with the vector of claim 18.
- 21. A process for producing a polypeptide comprising culturing a host cell according to claim 19 under conditions suitable for expression of said polypeptide.
- 22. A process for producing a polypeptide comprising culturing a host cell according to claim 20 under condition suitable for expression of said polypeptide.

23. An isolated polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
```

- 24. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 39.
- 25. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 44.
- 26. An isolated polypeptide capable of generating antibodies having binding specificity for a second polypeptide having a sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
```

27. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO : 39.

- 28. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO: 44.
- 29. An isolated polypeptide having an amino acid sequence selected from the group consisting of:

```
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
```

- SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
- SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
- SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
- SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
- SEO ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
- SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
- SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
- SEQ ID NO:40 and SEQ ID NO:41 or fragments, analogs or derivatives thereof.
- 30. The isolated polypeptide of claim 29 having an amino acid sequence according to SEQ ID NO : 39.
- 31. An isolated polypeptide having an amino acid sequence according to SEQ ID NO : 44.
- 32. An isolated polypeptide according to any one of claims 29 to 31, wherein the N-terminal Met residue is deleted.
- 33. An isolated polypeptide according to any one of claims 29 to 30, wherein the secretory amino acid sequence is deleted.
- 34. A vaccine composition comprising a polypeptide according to any one of claims 23 to 31 and a pharmaceutically acceptable carrier, diluent or adjuvant.

35. A vaccine composition comprising a polypeptide according to claim 32 and a pharmaceutically acceptable carrier, diluent or adjuvant.

- 36. A vaccine composition comprising a polypeptide according to claim 33 and a pharmaceutically acceptable carrier, diluent or adjuvant.
- 37. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 34.
- 38. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 35.
- 39. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 36.
- 40. A method according to any one of claims 37 to 39, wherein said animal is a bovine.
- 41. A method according to any one of claims 37 to 39, wherein said animal is a human.

42. A method according to any one of claims 37 to 39, wherein said bacterial infection is selected from the group consisting of group A streptococcus and group B streptococcus.

- 43. A method according to claim 42, wherein said bacterial infection is group B streptococcus.
- 44. Use of a vaccine composition according to claim 34 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 45. Use of a vaccine composition according to any one of claims 35 to 36 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 46. Use of a vaccine composition according to any one claims 23 to 31 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
- 47. Use of a vaccine composition according to claim 32 for the manufacture of a vaccine for the therapeutic or

prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

48. Use of a vaccine composition according to claim 33 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

S G K	E P A	N R F S	W A K	N K L	L I N G	60
•	AACTCTAGCA T L A	GCAACTATCT A T I L	TATTTTTTGC F F A	AGTTCAATTC V Q F	ATAGGTCTTA I G L K	120
AACCAGATTA P D Y	CCCTGGAAAA P G K	ACCTACTTTA T Y F I	TTATCCTATT I L L	GACAGCATGG T A W	ACTTTGATGG T L M A	180
CATTAGTAAC L V T	TGCTTTAGTG A L V	GGATGGGATA G W D N	ATAGGTATGG R Y G	TTCCTTCTTG S F L	TCGTTATTAA S L L I	240
TATTATTATT L L F	CCAGCTTGGT Q L G	TCAAGCGCAG S S A G	GAACTTACCC T Y P	AATAGAATTG I E L	AGTCCTAAGT S P K F	300
TCTTTCAAAC F Q T	AATTCAACCA I Q P	TTTTTACCGA F L P M	TGACTTACTC T Y S	TGTTTCAGGA V S G	TTAAGAGAGA L R E T	360
CCATCTCGTT I S L	GACGGGAGAC T G D	GTTAACCATC V N H Q	AATGGAGAAT W R M	GCTAGTAATC L V I	TTTTTAGTAT F L V S	420
CATCGATGAT S M I	ACTTGCTCTT L A L	CTTATTTATC L I Y R	GTAAACAAGA K Q E	AGATTAATAG D	AAAGTATCTA	480
GTGATAGACT	AACAGTATGA	TATGGTATGT	CAAAGTATTT	AGGAGGAGAA	GATATGTCTA M S T	540
CTTTAACAAT L T I	AATTATTGCA I I A	ACATTAACTG T _L T A		TTTTTATATT F Y I	> ATGTATTTGG M Y L E	600
AGACGTTAGC T L A	CACCCAGTCA T Q S	AATATGACTG N M T G	GGAAGATTTT K I F	TAGTATGTCT S M S	AAAGAAGAGT K E E L	660
TGTCATATTT S Y L	ACCCGTTATT P V I	AAACTTTTTA K L F K		TGTATACAAC V Y N	GGCTTGATTG G L I G	720
GCCTATTCCT L F L	CCTTTATGGG L Y G	TTATATATTT L Y I S	CACAGAATCA Q N Q	AGAAATTGTA E I V	GCTGTTTTTT A V F L	780
TAATCAATGT I N V	ATTGCTAGTT L L V	GCTATTTATG A I Y G		AGTTGATAAA V D K	AAAATCTTAT K I L L	840
TAAAACAGGG K Q G	TGGTTTACCT G L P	ATATTAGCTC I L A L	TTTTAACATT L T F	CTTATTTTAA L F	TACTACTTAG	900
CCGTTCGATT	TAGTTGAACG	GCTTTTAGTA	ATCATTTTT	TCTCATAATA	CAGGTAGTTT	960
AAGTAATTTG	TCTTTAAAAA	TAGTATAATA	TAACTACGAA	TTCAAAGAGA	GGTGACTTTG	1020
ATTATGACTG M T E	AGAACTGGTT N W L	ACATACTAAA H T K	GATGGTTCAG D G S D	ATATTTATTA I Y Y	TCGTGTCGTT R V V	1080
GGTCAAGGTC	AACCGATTGT P I V	TTTTTTACAT F L H	GGCAATAGCT G N S L	TAAGTAGTCG S S R	CTATTTTGAT Y F D	1140
AAGCAAATAG K Q I A	CATATTTTC Y F S	TAAGTATTAC K Y Y	CAAGTTATTG Q V I V	TTATGGATAG M D S	TAGAGGGCAT R G H	1200
					TGACTTAAAG	

GATATCTTAG TTCA D I L V H			GTTATATTGG V I L V	TAGGCCATAG G H S	CGATGGTGCC D G A	1320
AATTTAGCTT TAG			CCAGGTATGG P G M V		TTTGCTTAAT L L N	1380
TCAGGGAACC TGAG S G N L T			TGGTGGGATA W W D I	TTCTTTTAGT L L V	AAGGATTGCC R I A	1440
TATAAATTCC TTCA Y K F L H			TTTCCGTATA F P Y M	TGAGGCAAAA R Q K	AGCTCAAGTT A Q V	1500
ATTTCGCTTA TGT			AGTCCAGCTG S P A D	ATTTACAGCA L Q H	TGTGTCAACT V S T	1560
CCTGTAATGG TTT			ATAATTAAGT I I K L	TAAATCATTC N H S	TAAGAAACTT K K L	1620
GCTTCTTATT TTCC A S Y F P			TCTTTAGTTG S L V G	GCTTTGGGCA F G H	TCACATTATT H I I	1680
AAGCAAGATT CCCA K Q D S H			GCAAAAAAGT A K K F		TACGTTGAAA T L K	1740
GGAGAAATTG TTGA G E I V E		AATTGAAAA N	AGTCAAATCA	CTGACTTCTG	TGATTAAAAT	1800
TGTATTTTT ATA	TCTGTTT TA	AGTGCTTAT	TATTGTTGAA	ATGATTCATT M I H L	TGAAACGAAC K R T	1860
TATTTCTGTT GAG	CAACTAA AO Q L K		TGGGCAATTA G Q L		ATCTTTTCTT L F L	1920
AATTATCCTT GTG	GGGGTTA TO	CGCTGTCTT A V L	ACCGACAACC P T T	GGATATGACT G Y D F	TTGTACTGAA V L N	1980
TGGACTTTTA CGT.		AAGCAAAAG S K R			GGTGTATCAA C I N	2040
CACTTTTAAT AAC	TTGTCAG GA L S G I		CTTAATCGAT L I D		GCATGGCTTT M A F	2100
TTATGGTAAA AAA Y G K K				GTGACTCGTT V T R F	TTTTACCCTA L P Y	2160
TCTTATTTCT GGT	CTGTCAT T	TATTAGTGT I S V		ATCATGAGCC I M S H		2220
TGCCAAAGCT AGT A K A S	GTTGATT AC	CTATTATTT Y Y L	GGTATTAATT V L I (GGTGCTAGTA G A S M	TGTATTTTCC Y F P	2280
TGTTATTTAT TGG				TTCGGAGATA F G D M		2340
TACTCGTATA AAA T R I K	TTAGGTG T	TGTTTCTTT V S F	TTTTGAATGG F E W	GGATGTGCGG G C A A	CCGCAGCATT A A F	2400
TATAATTATC GGT I I I G			TCTACCAGTT L P V			2460

C :		G			V				S	L		P			L L		S	F	E	2520
ATTA(TT F	TAC:	AGG G	GT F	TTGC A	TGC A	CGA E	GGG <i>I</i> G	ACT.	ACCI P	r aa K	AGA E		TG V	TGGT'	TGC. A	ATG W	2580
		CTI L					CCTA Y			TAT					AGG' G		TCTA'	TTT F	CTT F	2640
			TT L	AGG' G	TAG' S	TC Q	AAAT I	AAA' N		ACGT R	ΓΤΑ' Υ		A AA N		CCC P		AAGA(E		AGT V	2700
ATCA S :	ACT T		CT L	ACA. Q	AAC T		TGGT V	GAG S		TTTC L	M	GCG1 R	I		AGG' G			CTT. L	AAT I	2760
ATTT:			A GC A	ATT' F	TTT' F	TG E	AAAA N	TAT'	TAC T	TTAT Y		TATO		GTT L	GCA Q		AGCT	AGG G		2820
	CCA P		A CA Q		ACA. Q			ATG W				AGG1 G			GCT L		GGGT'	ГТG' С	TTT F	2880
			A GC A	TAG R	AAC' T	TA I	TTGA D		AAA K	AGT(V	GAA. K		GC A	TTT F	TCC: P	AA I	TTGC'	TAT I	TAT I	2940
			TT L	GAC. T			TTTA Y		TAA N				I AT				GACTA L	ATC' S	TTT F	3000
		ATI I		ACT. L			TAGG G					raac K			TCT:		ATAA K		ACA Q	3060
ATTTA F			'AG S	CTG W	GGA. E	AG E	AGCG R		TAA K	GGA!	TGG: G				CGT' V		GTTTI L	AAT M	GGG G	3120
	I CTA	Y .TTI	S	W	E TGC.	E AG	R	I ACT.	K	D	G TAT	I	I G GC	I	V	S	L	M TGG	G	3120 3180
F CAGTTO	I CTA L GAA E	Y TTI F .CGC R	S TA Y CT	W TAT I GCA H	E TGC. A TTA Y	E AG G TA I	R GACT L TCAT I	I ACT L AGC A	K ATT F ATG W	D CCC: P GGA0 E	G TAT(I GCC(P	I CAGO R GATA I	I G GC A A GC A	I TCA H ATT L	V TAT I GGC A	TA T TA T	L CAGG G CGTT L	M TGG G GAT	G TAG S ICT L	
F C AGTTC V I I	I CTA L GAA E CTC L	Y TTT F .CGC R .GTT	S Y CT L TA	TATI I GCA H TTT	E TGC. A TTA Y ATG	E AG G TA I TT L	R GACT L TCAT I TGGT V	I ACT. L AGC. A TAA K	K ATT F ATG W GAT	D CCCT P GGAG E TTTI	G I GCC P ACA	I CAGO R GATA I AGGA	I G GC A GC A A A K	I TCA H ATT L ATC	V TAT' I GGC' A TTG'	TA TA TA TC Q	CAGGTC CGTTC L AGATT	M TGG G GAT I G G	G TAG S TCT L TGA D	3180
AGTTO V I	I CTA L GAA E CTC L ITC	Y TTT CGC R GTT V	S TA Y CT L TA Y OT V	W TAT I GCA H TTT L	TGC. A TTA Y ATG C TCG	AG G TA I TT L TT Y	R GACT L TCAT I TGGT V ATAA K	I ACT AGC A TAA K AAA K	K ATT F ATG W GAT I ACT L	CCCTP GGACE TTTT L ACTT	G TATO I GCCO P ACAL Q TCAL	I CAGO R GATA I AGGA AGCA	I G GC A GC A AA K K T TA	TCA H ATT L ATC S CGG	V TAT' GGC' A TTG' C TGG'	TA TA TC Q	CAGG	M TGG G GAT I GGA D	TAG S TCT L TGA D	3180 3240
AGTTO V I TACTO T I I TGTGT V I CGGTT G I	I CTA L GAA E CTC L ITC F ITA	Y TTI F CGCC R GTI V AAI N GCC	S TA Y CT L TA Y CT TT	W TAT I GCA H TTT L GGA TTT L	TGCA TTTA Y ATGC TCG R AAAA N	E AG G TA I TT L TT Y	R GACT L TCAT I TGGT V ATAA K ATAA	I ACT L AGC A TAA K AAA K	K ATT F ATG W GAT I ACT L GCT L	CCCCP GGAGE TTTTAL ACTTL CTAGY	G TAT I GCCC P ACA Q TCA Q CTG	CAGO R GATA G AGGA AGCA A GTAO	I G GC A GC A A A K T T A Y C C A	I TCA H ATT L ATC S CGG G AAA K	V TAT I GGC A TTG C TGG G AAA	TA TA TC Q TT S TG	CAGGE CGTTC AGAT I CTTCC S GAGAL E	M IGG G G G T G G G G A D A D	G TAG S TCT L TGA D TAG S TTG C	3180 3240 3300 3360 3420
AGTTO V I TACTO T I I TGTGTO V I CGGTTO V I	I CTA GAA E CTC TTC F TTA L GCG	Y TTTT F CGC R GTT V AAT N GCC A	S Y Y CT L TA Y GT V TT C CA C Q	W TATT I GCA H TTTT L GGGA D TTTT L ATT	TGC. A Y ATG C TCG R AAAA N TGT V	AG G TA I TT L TT Y TG D AA I	R GACT L TCAT I TGGT V ATAA K ATAA K TTGT V	I ACT. L AGC. A TAA K AAA K AAA N	K ATT F ATG W GAT I ACT L GCT L TAA	CCCCP GGAGE TTTTL L CTAGY TAAL	G TATC I GCCC P ACA Q TCA Q CTGC W ATG	CAGGATA GAGGAA GGTAG Y TCTT	I GGCA A GCCA A CTTA CTTA CTTA CTTA CTTA	I TCA H ATT L ATC S CGG G AAA K TAT	V TAT I GGCC A TTGG G A AAA N GGGG G	TA T TCQ TTS TGG GGE	CAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG	M TGG G G TIGG G G AGA D AGA AGC A	G TAG S TCT L TGA D TAG S TTG C CGG G	3180 3240 3300 3360 3420 3480
AGTTO V I TATTO T I I I I I I I I I I I I I I I I I	I CTA L GAA E CTC TTC F ITTA GCG A	Y TTTT F CGC R GTT V AAT N GCC A TTC T	S Y Y CT L TA Y OT T CT T T T T T T T T T T T T T T T T	W TATT I GCA H TTT L GGA D TTT L ATT F	E TGC A TTA Y ATG C TCG R AAAA N TGT V TCG R	AG G TA I TT Y TG D AA I TG E	R GACT L TCAT I TGGT V ATAA K ATAA K TTGT V AAGC	I ACT. AGC. A TAA K AAA K AAAG R CAA N TAT I	K ATT F ATG ATG ATG ACT L GCT L TAA N TGA	CCCCP GGACE TTTTA L ACTCL CTAC Y TAAAA K ATCC	G TATO I GCCO P ACA Q TCA Q CTGG W ATG C GTT F	CAGGAR GATA AAGGAA GTAG Y TCTT L TATT	G GC A GC A AAA AAA K T TAA C CAA C TAT I GAA D	I TCA H ATT L ATC S CGG G AAA K TAT M TGA D	V TAT' I GGC' A TTGG G AAAA' N GGG G TTGC	TAT TCQ TTS TGG GGE TGD	CAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG	M IGG G G AT I G G G AGA D AGC A G G C I G G C I G G C I G G C I G G C I G G C I G G C I G G G G	TAG S TCT L TGA D TAG S TTG C CGG G AGA D	3180 3240 3300 3360 3420 3480
TATTO TATTO TATTO TACTO TO TGGT CGGT CGTT C TGATO TGAT	I CTA L GAA CTC L CTC F GCG GAC GAC D	Y TTTT F CGCC R GTT V AAT N GCC TT CTT CTT	S Y Y CT L TA Y CT TA Y CT TA Y TA	W TATT I GCA H TTT L GGA D TTT T TAT I TTT F	E TGC A TTA Y ATG C TCG AAA TGT TCG R TTA	E AGG TA I TT Y TGD AA I TGE CA S	R GACT L TCAT I TGGT V ATAA K ATAA K TTGT V AAGC A GTAT I	AGCTAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	K ATT F ATG ATG ACT L GCT L TAA N TGA E ACA Q	GGACE TTTAL ACTTL CTACY TAAA K ATCC S GAACK	G TTATO	I CAGO R R GATA I AGGA A GTAO Y TCTT I GACA	I GAA CT	I TCA H ATT L ATC S CGG G AAA K TAT M TGA D ACT L	V TAT I GGC A TTG G G AAA N GGG G TTGC A TTGC A TTGC A	TA T TCQ TTS TGG GGE TGD ACH	CAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG	M IGG G GAT I IGG G G AGA D AGC A G G G G T L G G T G T G T G T G T G T G	TAG S TGA D TAG S TGG C G AGA D TGG G	3180 3240 3300 3360 3420 3480

	GAA	TAA	3 T.	ACAA	ACC'	ΤT	TCAG	AAA	TGC	CCT	AAA'	TAGA	GT'	TGA.	AAA	GG	ATGG	TTT	CTA	3720
G	N	K	Y	K	P	F	R	N	A	L	N	R	V	E	K	D	G	F	Y	
TTT F	'CGA E	AGT' V	r G V	TACA Q	ATC S	GC P	CACA H			AGA E					r r t L		AAGA E	GAT I		3780
TAA N	TAC T	TTG(W	G T'	TAGA E	AGG. G	AC R	GTCC P		AAA K			CTCA S			ATA' Y		TTAA N	TAA K		3840
TTA Y	TTT F	CCA? Q		AAGC A			TAGC A			AAA K		rgct A			CGA E			TGC A	TTT F	3900
TGC A	TAA N	TAT	r A' M				ATGA E			TAT I		CTCT ·s			rtt: L		TGCG R	TCA H	CGA D	3960
TAA K	ACA Q	GAA! K	A A				GCGT V					CTTT F		ATCI S		AT F	TCTC S	TTA Y		4020
TCA Q	AGA E	GAA(K	G G	GATA Y	CCA H	CT Y	ATTT F	TGA D	TTT L	GGG G	GAT(GGCA A	CC'	rtt <i>i</i> L			GAGT V	TGG G		4080
CGI V	TGA E	AACA T			rgc' A		AAGA E			GGC A							TCGG G	TAG S	TCA H	4140
TTT F	CTA Y	CTC! S	A T'	TTAA' N	TGG' G		TACA H		GTA Y			GAAG K		TACA T		AT L	TGTG W	GTC S	GGA E	4200
ACG R	TTA Y	TAT:	r T	CTTG	TTC' S	TC R	GTTC S			GTT.			GC: A					ATT. L	AAT M	4260
GGA E	AGA D	TAG:	r A. K		TAA K		TTGT V	TAA K	ATA	AGC	TTT	TTTA	' GG	CAA?	ATT	AA	AAGA	GCA	TGT	4320
CAI	GCG	ACA:	r G	CTCT	TTT	TA	AATC	ATT	TAA	TAC	CAT'	rga i	TG	CTT	GAA!	ГC	TACT	TTA	TAA	4380
TAT	'GAT	GTG	C T'	TTTA.	AAT.	ΑT	TGTT	TAG	CTA	CTG	TAG	CTGC	TG	ATT:	TAT	ЭC	ATTT	CAG	CTA	4440
CTI	GGT	AGT:	C.	ATTT	CTT	GC	ATTT	CTT	TTT	CAG	TGA'	TATG	AC	CAG	CAAC	ЭT	TTAT	TGA	GAG	4500
CTT	TTT	TTAG	C T'	ΓGA	(S	EQ	ID N	0:1)											4514

FIG. 1a [clonel-dna/aa]

SGKEPANRFS	WAKNKLLING	FIATLAATIL	FFAVQFIGLK	PDYPGKTYFI	50
	LVTALVGWDN				100
	TYSVSGLRET				150
KQED (SEQ					154
		FIG. 1	b		134
			~		
MSTLTIIIAT	LTALEHFYIM	YLETLATQSN	MTGKIFSMSK	EELSYLPVIK	50
	LIGLFLLYGL				100
	LALLTFLF				118
		FIG. 1	С		
	GSDIYYRVVG				50
VIVMDSRGHG	KSHAKLNTIS	FRQIAVDLKD	ILVHLEIDKV	ILVGHSDGAN	100
LALVFQTMFP	GMVRGLLLNS	GNLTIHGQRW	WDILLVRIAY	KFLHYLGKLF	150
PYMRQKAQVI	SLMLEDLKIS	PADLQHVSTP	VMVLVGNKDI	IKLNHSKKLA	200
SYFPRGEFYS	LVGFGHHIIK	QDSHVFNIIA	KKFINDTLKG	EIVEKAN	247
(SEQ ID NO:	: 4)				

FIG. 1d

MIHLKRTISV	EQLKSVFGQL	SPMNLFLIIL	VGVIAVLPTT	GYDFVLNGLL	50
RTDKSKRYIL	QTSWCINTFN	NLSGFGGLID	IGLRMAFYGK	KGQEKSDLRE	100
VTRFLPYLIS	GLSFISVIAL	IMSHIFHAKA	SVDYYYLVLI	GASMYFPVIY	150
WISGHKGSHY	FGDMPSSTRI	KLGVVSFFEW	GCAAAAFIII	GYLMGIHLPV	200
YKILPLFCIG	CAVGIVSLIP	GGLGSFELVL	FTGFAAEGLP	KETVVAWLLL	250
YRLAYYIIPF	FAGIYFFIHY	LGSQINQRYE	NVPKELVSTV	LQTMVSHLMR	300
ILGAFLIFST	AFFENITYIM	WLQKLGLDPL	QEQMLWQFPG	LLLGVCFILL	350
ARTIDQKVKN	AFPIAIIWIT	LTLFYLNLGH	ISWRLSFWFI	LLLLGLLVIK	400
PTLYKKQFIY	SWEERIKDGI	IIVSLMGVLF	YIAGLLFPIR	AHITGGSIER	450
LHYIIAWEPI	ALATLILTLV	YLCLVKILQG	KSCQIGDVFN	VDRYKKLLQA	500
YGGSSDSGLA	FLNDKRLYWY	QKNGEDCVAF	QFVIVNNKCL	IMGEPAGDDT	550
YIREAIESFI	DDADKLDYDL	VFYSIGQKLT	LLLHEYGFDF	MKVGEDALVN	600
LETFTLKGNK	YKPFRNALNR	VEKDGFYFEV	VQSPHSQELL	NSLEEISNTW	650
LEGRPEKGFS	${\tt LGYFNKDYFQ}$	QAPIALVKNA	EHEVVAFANI	MPNYEKSIIS	700
IDLMRHDKQK	IPNGVMDFLF	LSLFSYYQEK	GYHYFDLGMA	PLSGVGRVET	750
SFAKERMAYL	VYHFGSHFYS	FNGLHKYKKK	FTPLWSERYI	SCSRSSWLIC	800
AICALLMEDS	KIKIVK (SE	EQ ID NO:5)			816
		FIG.	1e		
MRILGAFLIF	STAFFENITY	IMWLQKLGLD	PLQEQMLWQF	PGLLLGVCFI	50
LLARTIDQKV	KNAFPIAIIW	ITLTLFYLNL	GHISWRLSFW	FILLLLGLLV	100
IKPTLYKKQF	IYSWEERIKD	GIIIVSLMGV	LFYIAGLLFP	IRAHITGGSI	150
ERLHYIIAWE	PIALATLILT	TAATCTAKIT	QGKSCQIGDV	FNVDRYKKLL	200
QAYGGSSDSG	LAFLNDKRLY	WYQKNGEDCV	AFQFVIVNNK	CLIMGEPAGD	250
DTYIREAIES	FIDDADKLDY	DLVFYSIGQK	LTLLLHEYGF	DFMKVGEDAL	300
VNLETFTLKG	NKYKPFRNAL	NRVEKDGFYF	EVVQSPHSQE	LLNSLEEISN	350
TWLEGRPEKG	FSLGYFNKDY	FQQAPIALVK	NAEHEVVAFA	NIMPNYEKSI	400
ISIDLMRHDK	QKIPNGVMDF	LFLSLFSYYQ	EKGYHYFDLG	MAPLSGVGRV	450
ETSFAKERMA	YLVYHFGSHF	YSFNGLHKYK	KKFTPLWSER	YISCSRSSWL	500

FIG. 1f

518

ICAICALLME DSKIKIVK (SEQ ID NO:6)

	GAAACAAC AACTTTTGAG E T T T F E	GCAATGAAAA A M K K			60
AAAATATCTG TT K I S V	GAGCGTTC TTTTATTGAA E R S F I E	TTTGATAAAC F D K L	TTCTATTAGC L L A	ACCTTATTGG P Y W	120
CGTAAAGGAA TG R K G M	CTGGCACT AATAGATAGT L A L I D S	CATGCTTTTA H A F N		ATGCTTAAAA C L K	180
	CAATTAAG CGCCTTTTTG Q L S A F L	TCCCAGTTAG S Q L D	ATAAAGATTT K D F	TTTATTTGAG L F E	240
	GCTTGGGC ATCACTCATC A W A S L I	TTGAGTATGG L S M E	AAGTTGAACA V E H	CACAAAGACT T K T	300
TTTTTAAAAA AA' F L K K '	TGGAAGAC ATCAACTCAC W K T S T H	TTTCAAAAAG F Q K D	ATGTTGAGCA V E H	TATAGTGGAT I V D	360
	CGTGAACA AATGGGATTG R E Q M G L	GCTAAAGAAC A K E H		TTATGGAAAA Y G K	420
	CAAGCGGA AGGTATTCGC Q A E G I R	AAAGCAAGAG K A R G		TGATTTCGAA D F E	480
AAAATAGAAC AAG K I E Q	CTAGATAG TGAGTTAGCA L D S E L A	ATCCATGATA I H D R	GGCATGAGAT H E I	AGTTGTCAAT V V N	540
GGTGGCACCT TAX	ATCAAGAA ATTAGGAATA I K K L G I	AAACCTGGTC K P G P	CACAGATGGG Q M G	AGATATTATC D I I	600
TCTCAAATTG AA' S Q I E	TTAGCCAT TGTTTTAGGA L A I V L G	CAACTGATTA Q L I N	ATGAAGAAGA E E E	GGCTATTTTA A I L	660
	CAGTACTT GATGGATTAG Q Y L M D	AGAGGATTAT	ATGAGCGATT M S D F	TTTTAGTAGA L V D	720
TGGATTGACT AAG G L T K	GTCGGTTG GTGATAAGAC S V G D K T	GGTCTTTAGT V F S	AATGTTTCAT N V S F	TTATCATCCA I I H	780
TAGTTTAGAC CG	TATTGGGA TTATTGGTGT I G I I G V	CAATGGAACT N G T	GGAAAGACAA G K T T	CACTATTAGA L L D	840
TGTTATTTCG GGT	TGAATTAG GTTTTGATGG E L G F D G	TGATCGTTCC D R S	CCTTTTTCAT P F S S	CAGCTAATGA A N D	900
	TTATTTAA AACAAGAACC Y L K Q E P			CAATTTTGGA I L D	960
CACCGTACTT TC	TTCTGACT TAAGAGAGAT S D L R E M		AAAGAATATG K E Y E		1020
TAATCACTAC GAAN H Y E	AGAAAGTA AGCAATCACG ESKQSR	TCTAGAGAAA L E K	GTAATGGCAG V M A E	AAATGGATTC M D S	1080
TTTAGATGCT TGG	GTCTATTG AGAGCGAAGT S I E S E V	CAAAACAGTA K T V	TTATCCAAAT L S K L	TAGGTATTAC G I T	1140
TGATTTGCAG TTC D L Q L	GTCGGTTG GTGAATTATC S V G E L S	AGGAGGATTA G G L	CGAAGACGTG R R R V	TTCAATTAGC Q L A	1200

GCAAGTATTA	TTAAATGATG	CAGATTTATT	GCTCTTAGAC	GAACCTACTA	ACCACTTAGA	1260
Q V L	L N D A	D L L	L L D	E P T N	H L D	
TATTGACACT	ATTGCATGGT	TAACGAATTT	TTTGAAAAAT	AGTAAAAAGA	CAGTGCTTTT	1320
I D T	I A W L	T N F	L K N	S K K T	V L F	
TATAACTCAT	GATCGTTATT	TTCTAGACAA	TGTTGCAACA	CGTATTTTTG	AATTAGATAA	1380
I T H	D R Y F	L D N	V A T	R I F E	L D K	
GGCACAGATT	ACAGAATATC	AAGGCAATTA	TCAGGATTAT	GTCCGACTTC	GTGCAGAACA	1440
A Q I	T E Y Q	G N Y	Q D Y	V R L R	A E Q	
AGACGAGCGT	GATGCTGCTA	GTTTACATAA	AAAGAAACAG	CTTTATAAAC	AGGAACTAGC	1500
D E R	D A A S	L H K	K K Q	L Y K Q	E L A	
TTGGATGCGT	ACTCAGCCAC	AAGCTCGTGC	AACGAAACAA	CAGGCTCGTA	TTAATCGTTT	1560
W M R	T Q P Q	A R A	T K Q	Q A R I	N R F	
TCAAAATCTA	AAAAACGATT	TACACCAAAC	AAGCGATACA	AGCGATTTGG	AAATGACATT	1620
Q N L	K N D L	H Q T	S D T	S D L E	M T F	
TGAAACAAGT	CGAATTGGGA	AAAAGGTTAT	TAATTTTGAA	AATGTCTCTT	TTTCTTACCC	1680
E T S	R I G K	K V I	N F E	N V S F	S Y P	
AGATAAATCT	ATCTTGAAAG	ACTTTAATTT	GTTAATTCAA	AATAAAGACC	GTATTGGCAT	1740
D K S	I L K D	F N L	L I Q	N K D R	I G I	
CGTTGGAGAT	AATGGTGTTG	GAAAGTCAAC	CTTACTTAAT	TTAATTGTTC	AAGATTTACA	1800
V G D	N G V G	K S T	L L N	L I V Q	D L Q	
GCCGGATTCG	GGTAATGTCT	CTATTGGTGA	AACGATACGT	GTAGGTTACT	TTTCACAACA	1860
P D S	G N V S	I G E	T I R	V G Y F	S Q Q	
ACTTCATAAT	ATGGATGGCT	CAAAACGTGT	TATTAATTAT	TTGCAAGAGG	TTGCAGATGA	1920
L H N	M D G S	K R V	I N Y	L Q E V	A D E	
GGTTAAAACT	AGTGTCGGTA	CAACAAGTGT	GACAGAACTA	TTGGAACAAT	TTCTCTTTCC	1980
V K T	S V G T	T S V	T E L	L E Q F	L F P	
ACGTTCGACA	CATGGAACAC	AAATTGCAAA	ATTATCAGGT	GGTGAGAAAA	AAAGACTTTA	2040
R S T	H G T Q	I A K	L S G	G E K K	R L Y	
CCTTTTAAAA	ATCCTGATTG	AAAAGCCTAA	TGTGTTACTA	CTTGATGAGC	CGACAAATGA	2100
L L K	I L I E	K P N	V L L	L D E P	T N D	
CTTAGATATT	GCTACATTAA	CTGTTCTTGA	AAATTTTTTA	CAAGGCTTTG	GTGGTCCTGT	2160
L D I	A T L T	V L E	N F L	Q G F G	G P V	
GATTACAGTT	AGTCACGATC	GTTACTTTTT	AGATAAAGTG	GCTAATAAAA	TTATTGCGTT	2220
I T V	S H D R	Y F L	D K V	A N K I	I A F	
TGAAGATAAC	GATATCCGTG	AATTTTTTGG	TAATTATACT	GATTATTTAG	ATGAAAAAGC	2280
E D N	D I R E	F F G	N Y T	D Y L D	E K A	
ATTTAATGAG F N E	CAAAATAATG Q N N E	AAGTTATCAG V I S	TAAAAAAGAG K K E	AGTACCAAGA S T K T		2340
AAAGCAAAGT	CGTAAAAGAA	TGTCTTACTT	TGAAAAACAA	GAATGGGCGA	CAATTGAAGA	2400
K Q S	R K R M	S Y F	E K Q	E W A T	I E D	
CGATATTATG	ATATTGGAAA	ATACTATCAC	TCGTATAGAA	AATGATATGC	AAACATGTGG	2460

D	I	M	I	L	Ε	N	T	I	T	R	I	E	:	N	D	М	Q	Т	С	G	
TA S		ATTTI F		CAA R						AA K				GAT D		AAA K	AA N	ATG.			2520
		AAAAG K	Y Y			GTT Y			ACCT L	TA(AGT L		GAC D	T	4		I	CCG' R		2580
AT' I		raaaa K N		TGA' D	TGA D			AGT' V		CAZ Q					AA	, AGT		ACG	CGC(CTAT Y	2640
GA' D	TTTA L	AGATA D K		ACC' P	I'GA' D	TAC T	AGC A		TTCA S		CCC P		L L	TAG D			TT L	GAC T	CTC S	ATAC Y	2700
TA Y	CGA) E	AAAAA K I		AGA(E	GAA K			ATT F		GT(V				AGA R			GA E	GAT'	TAT' I	TGGC G	2760
TG' C	TGG(G	CGGCT G F		TGG' G	rcc P		GAA K				TGC. A		GA M	TGC Q			GT V	GTA Y	CAT'	TGCA A	2820
GA. E	ACG' R	rttcc F R	: G'	TGG' G	raa K	GGG G		TGC' A				AGT V		AAA M			GA E	AGT.	AGA. E	AGCT A	2880
	AAA) K	AATTG I G		GTA' Y			ACT'		TTTA L	GA(GAC. T	AGC A	CA S	GTA T			AG S	TAG R	GGC A	AACT T	2940
		TTATA Y K		GCA' H	TAT M	GGG G	ATA' Y	TTG' C	TGCC A	TTZ L	ATC S	GCA Q	AC P				AA N	TGA' D		AGGT G	3000
CA H		AGCTA A M		GGA' D	TAT I		GAT(GA:		ATA	AG	TTG	AAA	AGT	GG	ATT	AGT	GAAC	3060
ΑT	GGA:	TAAT	T	ATT'	TTG.	AGA	TAA	GAG	GAAA	GA	AAA	.GGA	ιGΑ	CAT	ľ	4	GC A	Y	TAT' I	TTGG W	3120
TC S	TTA' Y	ITTGA L K		AAG R	GTA Y			TTG W		TG(W		TGA D			TAC		.GC		GCT' L	TTTT F	3180
GT V	GAC(T	GGTTA V I	T	CCT.	AGG. G	AAT M	GCC P	CAC. T	AGCC A		AGC A			TGA I			AA N	TGG G	CGT' V	TACA T	3240
AA K		IGATO D F		GAC' T		AGT V		TCT L			GTT F		CA M	TGT F			TT F	TGT V		ACTA L	, 330ó
GG G	TAT: I	IATTO I G	; G(GCG' R		TAC T	GAT M	GGC' A	TTAC Y	GC A	ATC S	TAG S	TC R	GCT			AC T	AAC. T	AAT M	GATT I	3360
AG. R	AGA: D	IATGO M F	: G'	TAA' N	TGA D	TAT M	GTA' Y	TGC A	TAAG K	CT' L	TCA Q	AGA E	TA. Y	ACT S	CCC	CAT H	CA H	TGA E			3420
CA Q	GAT I	AGGTG G V	T.	ATC' S		ACT L	AGT V			AT M					CT.				GAT M		3480
TT F	TGC' A	TGAAA E M	1 T			ACG R	TTT. L	AGG G	CCTA L	GT. V					TAZ		AT I	TTT F			3540
	TAT(M	GATAC I I		AAT I	TAC T	GAG S	TCC. P	ATC S	TTTG L	GC' A					CG(GCC P		3600
TT L	GGT. V	AGGAG G V	; T	CGT V	TTT. L	ATA Y	TGT. V	AGC A	TATA I	AA K	AAC T	AAA K	AAC P	CTT	TAT	rct s	GA E	AAG R	ACA O	ACAG	3660

ACTATGCTTG T M L D	ATAAAATCAA K I N	TCAATATGTT Q Y V	CGTGAAAATT R E N L		ACGCGTTGTT R V V	3 720
AGAGCCTTTG	CAAGAGAGAA	TTTTCAATCA	CAAAAATTTC	AAGTCGCTAA	CCAACGTTAC	3780
R A F A	R E N	F Q S	Q K F Q	V A N	Q R Y	
ACAGATACTT	CAACTGGTCT	TTTTAAATTA	ACAGGGCTAA	CAGAACCACT	TTTCGTTCAA	3840
T D T S	T G L	F K L	T G L T	E P L	F V Q	
ATTATTATTG	CAATGATTGT	GGCTATCGTT	TGGTTTGCTT	TGGATCCCTT	ACAAAGAGGT	3900
I I I A	M I V	A I V	W F A L	D P L	Q R G	
GCTATTAAAA	TAGGGGATTT	AGTTGCTTTT	ATCGAATATA	GCTTCCATGC	TCTCTTTTCA	3960
A I K I	G D L	V A F	I E Y S	F H A	L F S	
TTTTTGCTAT	TTGCCAATCT	TTTTACTATG	TATCCTCGTA	TGGTGGTATC	AAGCCATCGT	4020
F L L F	A N L	F T M	Y P R M	V V S	S H R	
ATTAGAGAGG	TGATGGATAT	GCCAATCTCT	ATCAATCCTA	ATGCCGAAGG	TGTTACGGAT	4080
I R E V	M D M	P I S	I N P N	A E G	V T D	
ACGAAACTTA	AAGGGCATTT	AGAATTTGAT	AATGTAACAT	TCGCTTATCC	AGGAGAAACA	4140
T K L K	G H L	E F D	N V T F	A Y P	G E T	
GAGAGTCCCG	TTTTGCATGA	TATTTCTTTT	AAAGCTAAGC	CTGGAGAAAC	AATTGCTTTT	4200
E S P V	L H D	I S F	K A K P	G E T	I A F	
ATTGGTTCAA	CAGGTTCAGG	AAAATCTTCT	CTTGTTAATT	TGATTCCACG	TTTTTATGAT	4260
I G S T	G S G	K S S	L V N L	I P R	F Y D	
GTGACACTTG	GAAAAATCTT	AGTAGATGGA	GTTGATGTAA	GAGATTATAA	CCTTAAATCA	4320
V T L G	K I L	V D G	V D V R	D Y N	L K S	
CTTCGCCAAA	AGATTGGATT	TATCCCCAA	AAAGCTCTTT	TATTTACAGG	GACAATAGGA	4380
L R Q K	I G F	I P Q	K A L L	F T G	T I G	
GAGAATTTAA	AATATGGAAA	AGCTGATGCT	ACTATTGATG	ATCTTAGACA	AGCGGTTGAT	4440
E N L K	Y G K	A D A	T I D D	L R Q	A V D	
ATTTCTCAAG	CTAAAGAGTT	TATTGAGAGT	CACCAAGAAG	CCTTTGAAAC	GCATTTAGCT	4500
I S Q A	K E F	I E S	H Q E A	F E T	H L A	
GAAGGTGGGA E G G S	GCAATCTTTC N L S		AAACAACGGT K Q R L	TATCTATTGC S I A	TAGGGCTGTT R A V	4560
GTTAAAGATC	CAGATTTATA	TATTTTTGAT	GATTCATTTT	CTGCTCTCGA	TTATAAGACA	4620
V K D P	D L Y	I F D	D S F S	A L D	Y K T	
GACGCTACTT D A T L	TAAGAGCGCG R A R	TCTAAAAGAA L K E	GTAACCGGTG V T G D		TTTGATAGTT L I V	4680
GCTCAAAGGG	TGGGTACGAT	TATGGATGCT	GATCAGATTA	TTGTCCTTGA	TGAAGGCGAA	4740
A Q R V	G T I	M D A	D Q I I	V L D	E G E	
ATTGTCGGTC	GTGGTACCCA	CGCTCAATTA	ATAGAAAATA	ATGCTATTTA	TCGTGAAATC	4800
I V G R	G T H	A Q L	I E N N	A I Y	R E I	
GCTGAGTCAC A E S Q	AACTGAAGAA L K N	CCAAAACTTA Q N L	TCAGAAGGAG S E G E	М	TGAGAAAAA R K K >	4860

ATCTGTTTTT S V F	TTGAGATTAT L R L W		AACTCGCTAC T R Y	AAAGCTACTC K A T L		4920
GATTTTTTG I F L	AAAGTTTTAT K V L S	CTAGTTTTAT S F M	GAGTGTTCTG S V L	GAGCCTTTTA E P F I	TTTTAGGGTT L G L	4980
AGCGATAACA A I T	GAGTTGACTG E L T A	CTAACCTTGT N L V	TGATATGGCT D M A	AAGGGAGTTT K G V S	CTGGGGCAGA G A E	5040
ATTGAACGTT L N V	CCTTATATTG P Y I A			TTTTTCAGAG F F R G	GTGTTTTCTA V F Y	5100
	TCTTATGGCT S Y G S	CAAATT (SE	Q ID NO:7)			5126

FIG. 2a

NFDIETTTFE	AMKKHASLLE	KISVERSFIE	FDKLLLAPYW	RKGMLALIDS	50
HAFNYLPCLK	NRELQLSAFL	SQLDKDFLFE	TSEQAWASLI	LSMEVEHTKT	100
FLKKWKTSTH	FQKDVEHIVD	VYRIREQMGL	AKEHLYRYGK	TIIKQAEGIR	150
KARGLMVDFE	KIEQLDSELA	IHDRHEIVVN	GGTLIKKLGI	KPGPQMGDII	200
SQIELAIVLG	QLINEEEAIL	HFVKQYLMD	(SEQ ID NO	:8)	229
		77.0			
		FIG. 2	b		
MSDFLVDGLT	KSVGDKTVFS	NVSFIIHSLD	RIGIIGVNGT	GKTTLLDVIS	50
GELGFDGDRS	PFSSANDYKI	AYLKQEPDFD	DSQTILDTVL	SSDLREMALI	100
KEYELLLNHY	EESKQSRLEK	VMAEMDSLDA	WSIESEVKTV	LSKLGITDLQ	150
LSVGELSGGL	RRRVQLAQVL	LNDADLLLLD	EPTNHLDIDT	IAWLTNFLKN	200
SKKTVLFITH	DRYFLDNVAT	RIFELDKAQI	TEYQGNYQDY	VRLRAEQDER	250
DAASLHKKKQ	LYKQELAWMR	TQPQARATKQ	QARINRFQNL	KNDLHQTSDT	300
SDLEMTFETS	RIGKKVINFE	NVSFSYPDKS	ILKDFNLLIQ	NKDRIGIVGD	350
NGVGKSTLLN	LIVQDLQPDS	GNVSIGETIR	VGYFSQQLHN	MDGSKRVINY	400
LQEVADEVKT	SVGTTSVTEL	LEQFLFPRST	HGTQIAKLSG	GEKKRLYLLK	450
ILIEKPNVLL	LDEPTNDLDI	ATLTVLENFL	QGFGGPVITV	SHDRYFLDKV	500
ANKIIAFEDN	DIREFFGNYT	DYLDEKAFNE	QNNEVISKKE	STKTSREKQS	550
RKRMSYFEKQ	EWATIEDDIM	ILENTITRIE	NDMQTCGSDF	TRLSDLQKEL	600
DAKNEALLEK	YDRYEYLSEL	DT (SEQ II	NO:9)		622
		FIG. 2	C		
		116. 2	C		
MIIRPIIKND	DQAVAQLIRQ	SLRAYDLDKP	DTAYSDPHLD	HLTSYYEKIE	50
KSGFFVIEER	DEIIGCGGFG	PLKNLIAEMQ	KVYIAERFRG	KGLATDLVKM	100
IEVEARKIGY	RQLYLETAST	LSRATAVYKH	MGYCALSQPI	ANDQGHTAMD	150

FIG. 2d

157

IWMIKDL (SEQ ID NO:10)

MAYIWSYLKR	YPNWLWLDLL	GAMLFVTVIL	GMPTALAGMI	DNGVTKGDRT	50
GVYLWTFIMF	IFVVLGIIGR	ITMAYASSRL	TTTMIRDMRN	DMYAKLQEYS	100
HHEYEQIGVS	SLVTRMTSDT	FVLMQFAEMS	LRLGLVTPMV	MIFSVVMILI	150
TSPSLAWLVA	VAMPLLVGVV	LYVAIKTKPL	SERQQTMLDK	INQYVRENLT	200
GLRVVRAFAR	ENFQSQKFQV	ANQRYTDTST	GLFKLTGLTE	PLFVQIIIAM	250
IVAIVWFALD	PLQRGAIKIG	DLVAFIEYSF	HALFSFLLFA	NLFTMYPRMV	300
VSSHRIREVM	DMPISINPNA	EGVTDTKLKG	HLEFDNVTFA	YPGETESPVL	350
HDISFKAKPG	ETIAFIGSTG	SGKSSLVNLI	PRFYDVTLGK	ILVDGVDVRD	400
YNLKSLRQKI	GFIPQKALLF	TGTIGENLKY	GKADATIDDL	RQAVDISQAK	450
EFIESHQEAF	ETHLAEGGSN	LSGGQKQRLS	IARAVVKDPD	LYIFDDSFSA	500
LDYKTDATLR	ARLKEVTGDS	TVLIVAQRVG	TIMDADQIIV	LDEGEIVGRG	550
THAQLIENNA	IYREIAESQL	KNQNLSEGE	(SEQ ID NO:	:11)	579
		FIG. 2	е		

MRKKSVFLRL	WSYLTRYKAT	LFLAIFLKVL	SSFMSVLEPF	ILGLAITELT	50
ANLVDMAKGV	SGAELNVPYI	AGILIIYFFR	GVFYELGSYG	SN	92
(SEQ ID NO:	:12)				

FIG. 2f

AATTTGGAAG F G S	TGCTCTATCA A L S	ACAGTTGAAG T V E V	TAAAGGAGAT K E I	TATTAGTGAA I S E	GAAAACATAT E N I W	60
	GCTCAGTTGC L S C	TGCCATTTTA C H F T	CTAGCTACTC S Y S	ATATTGGAAG Y W K	TTACCAACTT L P T W	120
GGTAAGCATC	ATATGGGTCT M G L	AGCAACAAAG A T K	GACAATCAGA D N Q I	TTGCCTATAT A Y I	TGATGACAGC D D S	180
AAAGGTAAGG K G K A		TAAAACAAAC K T N	AAAACGATGG K T M D	ATCAAATCAG Q I S	TGCTGAAGAA A E E	240
GGCATCTCTG G I S A		CGTAGTCAAA V V K	ATTACTGACC I T D Q	AAGGCTATGT G Y V	GACCTCACAC T S H	300
GGTGACCATT G D H Y	ATCATTTTTA H F Y	CAATGGGAAA N G K	GTTCCTTATG V P Y D	ATGCGATTAT A I I	TAGTGAAGAG S E E	360
TTGTTGATGA L L M T	D P N	TTACCGTTTT Y R F	AAACAATCAG K Q S D	ACGTTATCAA V I N	TGAAATCTTA E I L	420
GACGGTTACG D G Y V		CAATGGCAAC N G N	TATTATGTTT Y Y V Y		AGGTAGTAAG G S K	480
RKNI	R T K	ACAACAAATT Q Q I	A E Q V	A K G	T K E	540
AKEK	G L A	TCAAGTGGCC Q V A	H L S K	E E V	A A V	600
NEAK	R Q G	ACGCTATACT R Y T	T D D G	Y I F	S P T	660
DIID	D L G	AGATGCTTAT D A Y	LVPH	G N H	Y H Y	720
IPKK	DLS	TCCAAGTGAG P S E	L A A A	Q A Y	W S Q	780
KQGR	G A R	ACCGTCTGAT P S D	Y R P T	P A P	G R R	840
KAPI	P D V	GACGCCTAAC T P N	P G Q G	H Q P	D N G	900
GYHP	APP		D A S Q	N K H	Q R D	960
E F K G	KTF	KEL	L D Q L	H R L	TGATTTGAAA D L K	1020
TACCGTCATG Y R H V	TGGAAGAAGA E E D	TGGGTTGATT G L I	TTTGAACCGA F E P T	CTCAAGTGAT Q V I	CAAATCAAAC K S N	1080
AFGY	V V P		H Y H I	I P R	S Q L	1140
S P L E	M E L	AGCAGATCGA A D R	Y L A G	Q T E	D N D	1200
TCAGGTTCAG	AGCACTCAAA	ACCATCAGAT	AAAGAAGTGA	CACATACCTT	TCTTGGTCAT	1260

s	G	s	E	Н	s	K	P	s	D	K	E	v	Т	Н	·T	F	L	G	Н	
CC R	CA1 I	CAA K	AG A	CTTA Y	G G	AAA K	AGG G	CTT L	'AGAT D	GG G	TAA K	ACC P	AT Y	ATGA D	TAC	GAG S	TGF D	ATGC A	TTAT Y	1320
GT V	TTT F	TAG S	TA K	AAGA E	ATC S	CAT I	TCA H	TTC	AGTG V	GA D	TAA K	ATC S	AG G	GAGT V		AGC A	TAA K	ACA H	CGGA G	1380
G <i>P</i> D	ATCA H	TTT F	CC H	ACTA Y	TAT I	AGG G	ATT F	TGG G	AGAA E	CT L	TGA E	ACA Q	AT Y	ATGA E	GTT L	'GGA D	TGA E	GGT V		1440
A.F N	CTG W	GGT V	GA K	AAGC A	AAA K	AGG G	TCA Q	AGC A	TGAT D	GA E	GCT L	TGC A	TG A	CTGC A	TTT L			.GGA E	ACAA Q	1500
GG G	CAA K	AGA E	AA K	AACC P	ACT L	CTT F	TGA D	CAC T	TAAA K	AA. K	AGT V	GAG S	TC R		AGT V		AAA K		TGGT G	1560
AA K	AGT V	GGG G	CT Y	ATAT M	GAT M	GCC P	AAA K	AGA D	TGGT G	AA K	GGA D	CTA Y	TT F	TCTA'		TCG R		_	ACTT L	1620
GA D	TTT L	GAC T	TC Q	AGAT I	TGC A	CTT F	TGC A	CGA E	ACAA Q	GAZ E	ACT. L	AAT M	GC L	TTAA.	AGA D	TAA K	GAA K	GCA H	TTAC Y	1680
CG R	TTA Y	TGA D	CA I	TTGT V	TGA D	CAC T	AGG G	TAT I	TGAG E	CCZ P	ACG:	ACT L	TG A	CTGT: V		TGT V	GTC S	AAG S	TCTG L	1740
CC P	GAT M	GCA H	TG A	CTGG G	TAA' N	TGC A	TAC'	TTA Y	CGAT D	ACT	rgg: G	AAG S	TT S	CGTT'		TAT I	CCC.	ACA H	TATT I	1800
GA D	TCA H	TAT I	CC H	ATGT V	CGT' V	TCC P	GTA' Y	rtc. S	ATGG W	TTC L	GAC(GCG R	CG D	ATCA(GAT' I		AAC. T	AGT V	CAAG K	1860
TA Y	TGT V	GAT M	GC Q	AACA H	CCC(P	CGA E	AGT' V	rcg' R	TCCG P	GAT D	rgt <i>i</i> V	ATG: W	GT S	CTAA(GCA H	TGA. E	AGAG E	1920
TC S	AGG' G	TTC	GG V	TCAT'	TCC P	AAA N	TGT'	raco T	GCCT P	CTT L	rga: D	raa K	AC R	GTGCT		TAT M	GCC.	AAA N	CTGG W	1980
CA Q	AAT' I	TATO I	CC H	ATTC'	TGC: A	rga E	AGAZ E	AGT' V	ICAA Q	AA. K	AGC(A	CCT	AG A	CAGA <i>I</i> E	AGG' G	rcg R	TTT'	TGC A	AACA T	2040
CC. P	AGA(CGG(G	CT Y	ATAT:	TTT(F	CGA D	TCC2	ACG/ R	AGAT D	GT] V	TTTC L	GGC(CA K	AAGA <i>I</i> E	AAC: T	rtt F	TGT	ATG(GAAA K	2100
GA D	TGG(G	CTC(S	CT F	TTAG(CATO	CCC P	AAGA R	AGCA A	AGAT D	GGC G	CAGT S	rtc <i>i</i> S	AT L	TGAGA R	AAC(CAT I	TAA:	raaz K	ATCT S	2160
GA' D	rcta L	ATCO S	CC Q	AAGC: A	rgao E	GTG W	GCA <i>I</i>	ACAZ Q	AGCT A	CAA Q	GAC E	STT <i>I</i> L	AT L	TGGC <i>I</i> A	AA(K	GAA K	AAA: N	rac: T	rggt G	2220
GA' D	IGC: A	raci T	rg D	ATACO T	GGAT D	raa K	ACC(CAA! K	AGAA E	AAG K	CAA Q	ACA(Q	G A	CAGAT D	'AA(K	GAG S	CAA:			2280
CA.	ACA(Q	GCC <i>I</i> P	AA S	GTGAA E	AGCC A	CAG S	TAAA K	AGA <i>I</i> E	AGAA E	AAA K	GA <i>F</i> E	ATCA S	AG D	ATGAC D	TTT F	TAT I	AGA(2340
CCI P	AGA(D	CTAT Y	G G	GTCT?	AGA1 D	TAG R	AGCA A	ACC T	CCTA L	GAA E	GAT D	CAT H	A7 I	TCAAT N	CA <i>I</i> Q	ATT L	AGCA A			2400
GC' A	raat N	rato I	EG D	ATCC1	raac K	STA Y	TCTC L	EAT1	TTTC F	CAA Q	CCF P	AGA.A E	AG G	GTGTC V	CAF Q	ATT F	TTAT Y	raat N		2460

AATGGTGAAT TGGTAACTTA N G E L V T Y	TGATATCAAG A D I K T		AAATAAACCC I N P	TTAACCAAAA	2520
GAAGATCTCA TTGTTAAAGC	ACTGCTTTGT C	AAAGCAAGT	TACGGTGATT	TTGAAGTCAT	2580
TCTATGTAAC GAGTAGTGAT	AAAAGTTGGA T	AATAGCGGT	TTTCTTTTGC	AAAGAAATGG	2640
TATCCATGTT AGAATAGTAA	AAAAAGAGGA G	GATTCTTGG	ACTAATGTCA	AATAAGTAGA	2700
CAGAAAACTG TGTTATTTTA	TTGCGTTAAA A' I A N F	TAATTTTCT Y N E	TCTTTCTGAT E K Q	TAGGGGTTAG N P T L	2760
TCCTAGATTA GCCGTATGTG	GGTTGTAATT G	TTATAAAAA N Y F	TTCTCAATGT N E I	ATTCAAAGCA Y E F C	2820
GTCTAATTGA ACCTGTTTGA D L'Q V Q K	TATTTTGATA A' I N Q Y	TGTTTTCGG H K R	TTGATTTGTC N I Q	TATGCTTTAA R H K L	2880
ATACTTGAAA AATGCTTCAG Y K F F A E	TTACGGCATT A' T V A N	TCATAAGGA D Y P	TATCCAGGAT Y G P	TAGAAAAAGA N S F S	2940
ATGCATGATA TTGGCACTGC H M <	ACCCTAATAG TO	GAGACGCAA	GAAAAACACT	TTTAGGCAAT A I	3000
CAGTTTTCTG TACTGTACAG L K R Y Q V	GCGACTGGTC G' P S Q D	TTTAATCTC N L R	TGTTGAATTC Q Q I	TAGTTTCATT R T E N	3060
ATAAAATGTA ATGTAATTTT Y F T I Y N	TAACAATATT TO	GTTATACTA T I S	TCTTTGTTGT D K N	ATTTTCTCCT Y K R R	3120
ATTATGGAAA TAAAAGGTTT N H F Y F T		ACGGTGTGA V T H	AACCATTCAA F W E	TACAGGCATT I C A N	3180
ATCTGCAGGT GTTCCTTTTC D A P T G K	GAGACATTGA GO R S M S	CGGATAATG R I I	TCTTTTTCCG D K E	TGCAAGCCTG T C A Q	3240
GTAGTAAGCC ATAGAAGTAT Y Y A M <	ACACTGAGCC T	TGGTCACTG	TGTAAGATTG	CTCCTTTATT	3300
TAGGCAATTT TAACTGATTA K P L K L Q N		TACAAAATC V F D	CGTGTCCTGA T D Q	CAATCTGAGA C D S	3360
TAGTGTAAGC TATAATTTCT I T Y A I I E	CGGTTATAGA GA R N Y L		TGATGAGAGA S S L	TACAATTTAC Y L K	3420
AGTTACCGAA ATATAGGTAG C N G F Y L Y	GTAATATCTG TT	TACGAGCTT V L K	TTCCTTAGGC E K P	TTATCGGCAT K D A	3480
GGAAATCCCG ACTCAATTTA H G D R S L K	TTATCTGTTA AM	ATAATAAGC Y Y A	TTTACCCAAA K G L	TTGGGAACTT N P V	3540
TCTTGGTACG TGTCCGACAA K K T R T R C	AGCCAGCCAT TA	ATTTTTCAT N K M	GATACGATAG I R Y	ACTTTCTTTG V K K	3600
TATTAACAGT CAATCCGTGG	ATTTTTTTGA GO	CAATCGTGT . L R T	AATGGTACGA I T R	TAGCCATAAA Y G Y	3660
TAAAGTGATT CTCCATACAG I F H N E M <	AGCTGTTCAA T	TAATTCAAT .	AAGGTCATCT	TTTTTTGCGG	3720

CTTCTCATAC	TCCTTTTTC	AACGGTAATA	GGTCGACCGC	TTGACCTTAA	AACAGTCTAG	3780
AATGAAAACT	ATCGGGTAGT	TGTTTTTATA	GTCTTCCACA	AGCTTGATAA		3840
1 5 K	R I L G		KLL	DVQL	Q L Q	3900
TCCACTTCAG E V E	ACAGATGTTC S L H E	CAAGCCTTTA L G K	CCGTAGGTAT G Y T	ATTGCTTGCC Y Q K G		3960
TGAAAACGAT	AAAGCTCCTC	GTTTTCGTAC	CATTTCATCC	AAGTATAGAT	TTGACTATTA	4020
H F R	Y L E E	N E Y	W K M	W T Y I	Q S N	
TTTTTGATGC	CTAAAGTCTC	CATAATAACT	CTGTTAGACT	TGCCTGCTTT	CTTCATATCG	4080
N K I	G L T E	M I V	R N S	K G A K	K M D	
ATGCAAGCCA I C A	GCTTAGTTTC L K T E	CCATGAATAT W S Y	AKK	CCATAATAAA V M 	ACATTCCTGT	4140
TTCTAGTTTA	CTAAATTTCA	ACAGGAGTGT	TTTTCTTTTG	TCTCATTTTA	GGGATTCAGT	4200
GCCTATTGTT	GTCATCAATT	ATTTTTCTAA	ATTCCCCGGA	CTTAAATTGT	GACCCTTGGT	4260
CGGAATGAAA	GAGAAGTGTT	CCTTCAATCT	TTCTTTTATT	AAGTGAAAAG	GCAACACTTT	4320
TCTGTACAAC	ATTTATAAAG	TGTTTTTCTA	GGCAATTAAT . A I L	CTTTTAGTCA R K T	TTGGTGTTTG M P T Q	4380
GTAGTTGAGA	CTACCATGAA	TGCGGTGGTA	ATTCCACCAA	TGAACATAGT	CTTTAGTCTT	4440
Y N L	S G H	I R H Y	N W W	H V Y	D K T K	
AAGAGCTAGT	TCTTCCAGCA	ATTGAAAGGT	TTCTTGATAA	ACAAATTCAA	TTTTGAAAGC	4500
L A L	E E L	L Q F T	E Q Y	V F E	I K F A	
ACGATACGTA	CTTTCAGCTA	CGGCATTGTC	ATAAGGATAA	CCAGCCTGAC	TAAGCGAACG	4560
R Y T	S E A	V A N D	Y P Y	G A Q	S L S R	
TGTGATTCCA	AAGGCTTCCA	ATATTTCATC	AATTAACTGA	TTATCAAACT	CTTTGCCACG	4620
T I G	F A E	L I E D	I L Q	N D F	E K G R	
ATCTGAATGG	AACATCTTGA	CTTTGGTCAG	GGCGTAAGGG	ATGCTTTGTA	TGGCTTGCTT	4680
D S H	F M K	V K T L	A Y P	I S Q	I A Q K	
AACGAGTTCA	GCGGTCTTGT	GCCAACCAAG	AGACAGGCCG	ATGATTTCAC	GGTTGTATAG	4740
V L E	A T K	H W G L	S L G	I I E	R N Y L	
GTCAATGATG	AGGCAAACAT	AAGCCCAACG	ATTGCCTACA	CGAACATAGG	TTAAGTCAGT	4800
D I I	L C V	Y A W R	N G V	R V Y	T L D T	
GACTAAGGCT	TGTAGTGGTC	TTTCTTGCTT	AAATTGCCTG	TCTAAGTGGT	TGGGAATAGG	4860
V L A	Q L P	R E Q K	F Q R	D L H	N P I P	
A E N	K G K	AATGTGGTTT S H P K	FTA	K Q Y	V S V L	4920
NLK	V W I	GTCGAATCCG R R I R	RRS	L T I	GENN	4980
CAAGCATATT	TTGATTTTTC	TGGATCCGTA	TCTAGACTCG	CTATCGAGAA	AAATTCTTTT	5040
L C I	K I K	R S G Y	R S E	S D L	F I R K	

110 //142500				T C I / C/L	////
AATAGTTTCT TO				AGT AATAACTTGA Y Y S S	5100
GTGTGGCATA TI H P M N	FCAGCCAGC GACA N L W R C			CCT TATTAGCAGT	5160
GATTATTTCC CT I I E F (SEQ ID NO:	R K T G Y	AATCACC GCTGCT / D G S S FIG. 3	S A K P Y		5215
FGSALSTVEV (SEO ID NO:		LYRLSCCHFT	SYSYWKLPTW		40
(019 10 10.	111				
		FIG. 3	3b		
MGLATKDNQI	AYIDDSKGKA	KAPKTNKTMD	QISAEEGISA	EQIVVKITDO	50
		AIISEELLMT			100
		RTKQQIAEQV			150
		YIFSPTDIID			200
DLSPSELAAA	QAYWSQKQGR	GARPSDYRPT	PAPGRRKAPI	PDVTPNPGQG	250
HQPDNGGYHP	APPRPNDASQ	NKHQRDEFKG	KTFKELLDQL	HRLDLKYRHV	300
EEDGLIFEPT	QVIKSNAFGY	VVPHGDHYHI	IPRSQLSPLE	MELADRYLAG	350
QTEDNDSGSE	HSKPSDKEVT	HTFLGHRIKA	YGKGLDGKPY	DTSDAYVFSK	400
ESIHSVDKSG	VTAKHGDHFH	YIGFGELEQY	ELDEVANWVK	AKGQADELAA	450
ALDQEQGKEK	PLFDTKKVSR	KVTKDGKVGY	MMPKDGKDYF	YARDQLDLTQ	500
IAFAEQELML	KDKKHYRYDI	VDTGIEPRLA	VDVSSLPMHA	GNATYDTGSS	550
FVIPHIDHIH	VVPYSWLTRD	QIATVKYVMQ	HPEVRPDVWS	KPGHEESGSV	600
IPNVTPLDKR	AGMPNWQIIH	SAEEVQKALA	EGRFATPDGY	IFDPRDVLAK	650
ETFVWKDGSF	SIPRADGSSL	RTINKSDLSQ	AEWQQAQELL	AKKNTGDATD	700
TDKPKEKQQA	DKSNENQQPS	EASKEEKESD	DFIDSLPDYG	LDRATLEDHI	750
NQLAQKANID	PKYLIFQPEG	VQFYNKNGEL	VTYDIKTLQQ	INP	793

FIG. 3c

(SEQ ID NO:15)

MTDPNYRFKQ	SDVINEILDG	YVIKVNGNYY	VYLKPGSKRK	NIRTKQQIAE	50
QVAKGTKEAK	EKGLAQVAHL	SKEEVAAVNE	AKRQGRYTTD	DGYIFSPTDI	100
IDDLGDAYLV	PHGNHYHYIP	KKDLSPSELA	AAQAYWSQKQ	GRGARPSDYR	150
PTPAPGRRKA	PIPDVTPNPG	QGHQPDNGGY	HPAPPRPNDA	SQNKHQRDEF	200
KGKTFKELLD	QLHRLDLKYR	HVEEDGLIFE	PTQVIKSNAF	GYVVPHGDHY	250
HIIPRSQLSP	LEMELADRYL	AGQTEDNDSG	SEHSKPSDKE	VTHTFLGHRI	300
KAYGKGLDGK	PYDTSDAYVF	SKESIHSVDK	SGVTAKHGDH	FHYIGFGELE	350
QYELDEVANW	VKAKGQADEL	AAALDQEQGK	EKPLFDTKKV	SRKVTKDGKV	400
GYMMPKDGKD	YFYARDQLDL	TQIAFAEQEL	MLKDKKHYRY	DIVDTGIEPR	450
LAVDVSSLPM	HAGNATYDTG	SSFVIPHIDH	IHVVPYSWLT	RDQIATVKYV	500
MQHPEVRPDV	WSKPGHEESG	SVIPNVTPLD	KRAGMPNWQI	IHSAEEVQKA	550
LAEGRFATPD	GYIFDPRDVL	AKETFVWKDG	SFSIPRADGS	SLRTINKSDL	600
SQAEWQQAQE	LLAKKNTGDA	TDTDKPKEKQ	QADKSNENQQ	PSEASKEEKE	650
SDDFIDSLPD	YGLDRATLED	HINQLAQKAN	IDPKYLIFQP	EGVQFYNKNG	700
ELVTYDIKTL	QQINP (SEQ	Q ID NO:16)			715
		FIG. 3	d		
MHSFSNPGYP	YDNAVTEAFF	KYLKHROINR	KHYONIKOVO	LDCFEYTENE	50
	LGLTPNQKEE				77
		,		,	,,
		FIG. 3	е		
MAYYOACTEK	DITEMEDUC				
	DIIRSMSRKG				50
DOTINIVINI	ITFYNETRIQ	OKTUDÖSÞAÖ	YRKLIA (SE	Q ID NO:18)	86
		FIG. 3	f		
MENUETVOVO	TTTD	CI MINIMATERIA			
	TITRLLKKIH				50
	LSRDFHADKP				100
TIATTISDCQ	DTDFVLDTLN	QLKLPK (SE	Q ID NO:19)		126

FIG. 3g

MVKKAYSWET I	KLACIDMKKA	GKSNRVIMET	LGIKNNSQIY	TWMKWYENEE	50
LYRFHQGVGK (QYTYGKGLEH	LSEVEQLQLQ	VDLLKKYRGL	IRKSIK	96
(SEQ ID NO:2	20)				
		ETC 2	1		

FIG. 3h

IRYPKASSGDYGTKREIITANKDKYSISKMCRWLNMPHSSYYYQAVESVS50ETEFEETIKRIFLDSESRYGSRKIKICLNNEGITLSRRRIRRIMKRLNLV100SVYQKATFKPHSRGKNEAPIPNHLDRQFKQERPLQALVTDLTYVRVGNRW150AYVCLIIDLYNREIIGLSLGWHKTAELVKQAIQSIPYALTKVKMFHSDRG200KEFDNQLIDEILEAFGITRSLSQAGYPYDNAVAESTYRAFKIEFVYQETF250QLLEELALKTKDYVHWWNYHRIHGSLNYQTPMTKRLIA(SEQ ID NO:21) 288

FIG. 3i

AA	TTT	GAA	AG	CAGA	ATT.	ATC	TGT	AGA	AGAT	GA	GCA.	ATA	AT.	CAGC	AAC	AGT	TTA	TGG	TAAA	60
N	L >	K	A	E	L	S	V	E	D	E	Q	Y	Т	A	Т	V	Y		K	
TC S	TGC A	TCA H	TG G	GTTC.	AAC. T	ACC P	ACA Q	AGA E	AGGT G	GT V	TAA N	TGG G	GG A	CGAC T	TTA Y		AGC A		TTAT Y	120
CT. L	AAG S	TCA. Q	AT F	TTGA	TTT' F	TGA E	AGG G	TCC P	TGCT A	CG R	TGC A	TTT F	CT L	TAGA D	TGT V		AGC A		CATT I	180
AT' I	TCA: H	CGA. E	AG D	ACTT F	CTC S	AGG G	TGA E	AAA K	ACTT L	GG.	AGT. V	AGC A	TT Y	ATGA E	AGA D	TGA D	CTG C	TAT M	GGGA G	240
CC. P	ATT L	GAG(S	CA M	TGAA' N	TGC A	AGG G	TGT V	CTT F	CCAG Q	TT F	TGA' D	TGA E	AA T	CTAA' N	TGA' D	TGA D	TAA' N	TAC'	TATC I	300
GC' A	L L	raa' N	TT F	TCCG'	TTA(Y	CCC P	ACA.	AGG G	GACA T	GA D	TGC' A	TAA K	AA T	CTAT		AAC T	TAA			360
AA K	ACT' L	raa N	CG G	GAGT'	rgaz E	AAA K	AGT V	GAC' T	TCTT L	TC S	TGA D	CCA'	TG E		CAC T		ACA H		TGTA V	420
CC' P	rato M	GGA(D	CG D	ATGAZ E	ATTI L	AGT V	ATC:	AAC T	CTTA L	CT. L	AGC' A	TGT V	CT Y	ATGA E		GCA Q	AAC' T			480
AAZ K	AGG2 G	ACA' H	rg E	AACA(Q	GGT' V	TAT I	TGG' G	TGG' G	rggg G	AC. T	ATT' F	IGG G	TC R	GCTT.	ACT' L	rga E	ACG		TGTT V	540
GCZ A	ATA Y	CGG' G	rg A	CCAT(GTT(F	CCC P	AGG G	AGA' D	rgaa E	AA N	CAC' T	rat M	GC H		AGC' A			GTA Y		600
CC:	CTT	AGAZ	A.F	TATA	rtt	CCG	TTC	GGC'	rgct	AT	CTA	CGC.	AG	AAGC'	TAT(CTA	TGA	ATT	AATC	660

P	L	E	. N]	-	F	R	S	A	A	I	Y	A	E	А	I	Y	E	L	I	
AA K	AT <i>I</i>	AAA	ATA	. ATC	CT	TA	AAC	TAA	AT <i>I</i>	ATGTG	AT	CAF	ATG	ATA	AAGG	GT	GTG	AA	GACA	ATGAA	720
AG	TGT	CT	TTG	CCI	CT'	TT'	TCA	TAA	.GG1	TAGA	тт	TGG	SAGA	ACT	M	T	D	CT1 L		AAAA K	780
AT' I	rat I	TTA K	AAG A	CAA	TA	AA K	AAG S	TGA D	TTC S	CACAG Q	AA N	TCA Q		TTA Y	ATAC T		-> \AAA N	TG0 G	TAT I	TGAT D	840
CC' P	rti L	GT F	TTG A	CTC A	CT	CC! P	raa K	AAC T	AGC A	TAGG R	AT I	CAA N		TTG V		CC <i>I</i> Q	AAGC A	ACC P	CTGG G	STTTA L	900
AA K	AAC T	CTC Q	AAG E	AAG A	CA	AG2 R	ACT L	CTA Y	TTC W	GAAA K	GA D		ATC S	CTG G	GAGA D	TC(R	STCT L	ACC R	GCCA Q	.GTGG W	960
CT: L	rgo G	AG V	TTG D	ATG E	AA(GA(E	GAC T	ATT F	TTA Y	CCAT H	TC S	TGG G	AA <i>I</i> K	AT F		TGI V	TTT L	ACC P	TTT L	'AGAT D	1020
TT:	TTA Y	TTA Y	ACC P	CAG	GC2	AA <i>I</i> K	AGG G	AAA K	ATC S	AGGA G	GA D	TTT L		CCC P	CTAG R	AA. K	AGG G	TTT F	TGC A	GGAG E	1080
AA) K	OTA W	GC H	ACC P	CTC	TTZ	AT: I	TTT L	AAA K	AGA E	AATG M	CC'		TG1 V		AATT L	GAC T	CTT L	GCT L	'AGT V	TGGT G	1140
CA(Q	STA Y	TG A	CTC Q	AGA K	AA:	ra: Y	ATT Y	TCT L	TGG G	AAGC S			ACA H		AAAA N	TCI L	'AAC T	AGA E	AAC T	AGTT V	1200
AAA K	AGC A	TT. Y	ACA K	AAG D	AC'	rat Y	rct L	ACC P	CGA D	TTAT Y	TT. L	ACC P		'GG V	TTCA H	CCC P	ATC S	ACC P	GCG R	AAAT N	1260
CA/ Q	I	TT W	GGC L	TAA K	AGA I	AAC K	SAA N	TCC P		GTTT F	GAZ E	AAA K	AGA D	TC L	TAAT I	CGI V	TGA D		'ACA Q	AAAG K	1320
AT <i>I</i>	AGT V	'AG A	CAG D	ATA I	TTT I	rt <i>r</i>	AAA K	AGA D	TTA •	AGGA	TA	GGA	GTI	'GG	TATG	R	D			CTAC L H	1380
ACA	ACG	TA' Y	TTT F	TTC S	CT <i>I</i> Y	OTA 1	SAT	TGT C	CAA Q	ACGG T A	CA'	TTT F	GAG E	GA D	CTAT	ATT	AAT N			ACAG T G	1440
GT C	SAA :	TT' F	TAT I	CAC T	GA(T	CAG	SAA E	CAT H	TTT F	GATT D L		TCA S		CC P	TTAC.		GGT G			GATG D V	1500
TTC	CT	GA' D	TTA Y	TAG S	TGC A	TT Y	AT ?	TGT C	CAA Q	AAAA K I	TAG	GAT D	TAT Y	CT L	TAAT	CAG Q	AAA K	TAT Y	GGA G	AATC N R	1560
GAT E	TT	AA K	AAA K	AGG. G	IAA I	TC E	AA I	ATC I	GGT G	TATT Y F		AAA K		AG R		TCA S				GATT D Y	1620
ATT I	TA	AA K	AAA N	TAA K	AG <i>P</i> E	ra <i>l</i>	TT	GAT D	TTA L	AAAC K L	TA:			AT I			AAT N			TATG Y D	1680
TTA Y	'AT	L L	GCA Q	AGA. E	AGA E	AA A	CT Y	CTG.	AAA K	GTAC V P	CAZ	ACA. T		GG G			AGC S	AGA R		CTTT L .	1740
AAI	'CG	TA!	rgg	AAT	TTG	CC	TAC	AGG	CCG	TGTG	GA.	AGC	GCA	.CG	TTTT	AGC	TCA	CTT	TGA	TTAT	1800
GGI	'TT	TC	GTA	AGT'	TAP	AAC	TT	AGA	TGT.	AGAA	GA:	TTT.	AAA	AC	CGTT'	ГGА	AAC	GCA	ATT	GAAG	1860
CGC	ΑT	TT:	ГСА	TAA	AGA	ATG	TT	ATC'	TAA	GGGG	TTA	AGC'	TTT	TG	AACT	AAA	TAC	CAA	ATC	CCTT	1920

TATCTATATG	GGAATGAAAA	ACTTTATCG	C TATGCTTTAG	AGATACTCAA	ACAGCTTGGT	1980
			CATATTCCTG			2040
GATAGACTTC	AAGGTCTGCT	AAAGGACTA	CAAATTGATG	AAAATCATTT	GATATGAGGA	2100
AATTTTTGAT	AAAAAAGCTA	GGCAATATT	G CTTAGCTTTT	TTGTAATGCT	ATTGATAGTT	2160
TTAGTGAAAA	TTTCAAAAAA	ATAAAGAAA	CATTTACTTG	TTGCAAGCGC	TTGCGTAAAT	2220
			A AAAAAGGAGA	М К 	RKD	2280
TTATTTGGTG L F G D	ATAAACAAAC K Q T	TCAATACACO Q Y T	ATTAGAAAGT I R K L	TAAGTGTTGG S V G	AGTAGCTTCA V A S	2340
	v C 1	гьн	AGTCCACAGG S P Q V	FAE	E V S	2400
	I I A	I A L	TCGAATATTA S N I N	Q V D	и о о	2460
TCTACTAATT S T N L	TAAAAGATGA K D D	CATAAACTCA I N S	AACTCTGAGA N S E T	CGGTTGTGAC V V T	ACCCTCAGAT P S D	2520
	кΩг	v S D	GAAACTGACA E T D T	Q K G	V T E	2580
11	1 3 1	ד ב ב	AATAAAGGTC N K G P	V S D	KNT	2640
TTAGATTTAA L D L K	AAGTAGCACC V A P	ATCTACATTG S T L	CAAAATACTC Q N T P	CCGACAAAAC D K T	TTCTCAAGCT S Q A	2700
ATAGGTGCTC I G A P	CAAGCCCTAC S P T	CTTGAAAGTA L K V	GCTAATCAAG A N Q A	CTCCACGGAT P R I	TGAAAATGGT E N G	2760
TACTTTAGGC Y F R L	TACATCTTAA H L K	AGAATTGCCT E L P	CAAGGTCATC Q G H P	CTGTAGAAAG V E S	CACTGGACTT T G L	2820
TGGATATGGG W I W G	GAGATGTTGA D V D	TCAACCGTCT Q P S	AGTAATTGGC (CAAATGGTGC N G A	TATCCCTATG I P M	2880
ACTGATGCTA T D A K	AGAAAGATGA K D D	TTACGGTTAT Y G Y	TATGTTGATT T	TTAAATTATC K L S	TGAAAAACAA E K Q	2940
CGAAAACAAA R K Q I	TATCTTTTT . S F L	AATTAATAAC I N N	AAAGCAGGGA (CAAATTTAAG N L S	CGGCGATCAT G D H	3000
CATATTCCAT H I P L	TATTACGACC L R P	TGAGATGAAC E M N	CAAGTTTGGA C	TTGATGAAAA D E K	GTACGGTATA Y G I	3060
CATACTTATC . H T Y Q	AACCCCTCAA. P L K	AGAAGGGTAT E G Y	GTCCGTATTA A	ACTATTTGAG Y L S	ITCCTCTAGT S S S	3120
AACTATGACC . N Y D H	ACTTATCAGC A	ATGGCTCTTT W L F	AAAGATGTTG (CAACCCCYTC A	AACAACTTGG T T W	3180
CCAGATGGTA (P D G S	GTAATTTTGT (N F V	GAATCAAGGA N Q G	CTATATGGAA (GTATATTGA '	IGTATCACTA V S L	3240

K :	ACT <i>I</i>	AACG 1 A	CCAA K	AGA E	GAT I	TGG G	TTT F	TCTA L	ATO I	CTT. L	AGA1 D	G E	AAAG S	TA <i>F</i> K			SAGA D		3300
GTG? V F	AAAC K V	TTC VQ	AACC P	CAA N	CGA D	CTA Y	TGT V	TTTT F	AGA R	AGA' D	TTTA L	AG A	CTAA N	CCA H	ATAA N	CCA Q	LAAT I	TTTT F	3360
GTA.	AAAG K [ATA K	AGGA D	TCC P	AAA K	GGT V	TTA Y	TAAT N	AA1 N	rcc' P	TAT Y	T Y	ACAT I	TGA D	TCA Q		'GCA Q	GCTA L	3420
K) A	ı Q	~	Ι	D	L	Т	S	Ι	Q	A	S	F	Т	Т	L	D	G	3480
V E) K	Т	CTGA E	Ι	L	K	E	L	K	V	T	D	K	N	Q	N	A	I	3540
CAAA Q I	ATTT S	CTG D	ATAT	CACT T	rct L	CGA'	TAC T	TAGT S	AAA K	ATC: S	CTT L	T L	TAAT	AAT I	CAA K		CGA D	CTTT F	3600
AATC N F	CTA K	AAC Q	AAGG' G	TCAT H	TTT F	CAA(N	CAT: I	ATCT S	TAT Y	'AA' N	GGT G	A N	ACAA' N	TGT V	CAT M	GAC T			3660
TCTT S W	GGG E	AAT F	TTAA: K	AGAC D	CA Q	ACT:	ΓΤΑ' Υ	IGCT A	TAT Y	'AG' S	GGA.	A. N		AGG G	TGC A	AGT V		CAAT N	3720
CAAG Q D	ATG G	GTT S	CAAA K	AGTI V	GA E	AGC(CAG(S	CCTC L	TGG W	TC <i>I</i> S		A S	GTGC: A	rga D	TAG S	TGT V	CAC' T	TATG M	3780
ATTA I I	TTT. Y	ATG D	ACAA K	AGA'I D	'AA N	CCAA Q	AAA(N	CAGG R	GTT V	'GT <i>F</i> V	GCG. A	A (CCT L	TGT V		AAAT N	3840
AATA N K	AAG G	GTG V	TTTG(W	GCAG Q	AC T	GAT <i>I</i>	ACT: L	rgat D	ACT T	AAA K	L (G (G		raa. K		CTA' Y	TAC'	rggt G	3900
N K	ATT	V ATC	TTTG(W TTTA(Y	Q CGAA	Т	Ι	L AAGA	D AGGT	T AAG	K GAI	L	G G !	Ι	K SAT'	N TTT	Y	T	G	3900 3960
TACT Y Y	ATT Y AGT	ATC L CAT	W	Q CGAA E AGAG	T AT I	I AAAA K	L AAGI R TAGI	D AGGT G	T AAG K ACT	K GAI D GTI	L (AAG(K '	G G ' V G 1	I TTAAG K	K SAT' I	N ITT L IAA	Y AGA' D	T TCC: P GGC:	G ITAT Y IAAA	
TACT Y Y GCAA A K	ATT. AGTO	ATC L CAT L	TAGCA	Q EGAA E AGAG E	T AT I TG W	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R TAGT S	D AGGT G FAAT N	T AAG K ACT T CCT	K GAI D GTI V CAA	L (AAGO K 'AATO N I	G G V G D	I TTAAG K ATGAT D	K I I IAT' I	N ITT L IAA K	Y AGA' D AACO T	TCC: P GGC: A	G TTAT Y TAAA K	3960
TACT Y Y GCAA A K GCAG A A	ATT. AGT(S CTT' F	ATC L CAT L ITG V	TAGCA	Q EGAA AGAG E ICCA P	T IAT I TG W AG S	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R TAGT S ACTT L	AGGT G IAAT N IGGA	AAG K ACT T CCT P TAC	K GAI D GTI V CAA Q	L (AAGO K 'AATO N IAATO N IAAT	G G V G D D T T T C	I TTAAC K ATGAT D TAAGT	K I I I I I I I I I I I	N ITT L IAA K IGC A	Y AGA' D AACC T TAAA	TCCTPGGCTA	G ITAT Y IAAA K IGCT A	3960 4020
TACT Y Y GCAA A K GCAG A A AATT N F	AGTO AGTO F TTAI	V ATC L CAT L ITG V AAG G	TAGCA TAGAA TAAAA N	Q EAAGAG ECCA P ACAA Q	T AT I TG W AG S GA D	I AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R FAGT S ACTT L CGTT V	AGGT G IAAT N IGGA CATA I	AAG K ACT T CCT P TAC Y	K GAI GTI V CAA Q GAA E	AAGG K 'AATG N I AATG N I GCAG A I	G G Y G D T L CH	I TTAAG K ATGAT D TAAGT S ATGTA V	K SAT' I SAT' I TT' F	ITT LIAA KIGC AGA D	AGA' D AACC T TAAA K CTTC	TCCTP GGCTA AATTI CACTT	G ITAT Y IAAA K IGCT A ITCT S	3960 4020 4080
TACT. Y Y GCAA A K GCAG A A AATT. N F GATC. D R	AGTON S CTTT TTAL GATON TAGA	V ATC L CAT L FTG V AAG G CTT L ATT	TTTAC Y TAGCA A TAAAT N GAAGA R TGGAT	Q EGAA E CCA P ACAA Q GGA	T AT I STG W AG S GA D AA K AA	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R R S ACTT L CGTT V AAAA	AGGT G FAAT N FAAT N LAAT N LAAT N	AAG K ACT T CCT P TAC Y CAA	K GAI D GTI V CAA Q GAA E TTT F	AAAGGK NAATGAATGAATGAATGAATGAATGAATGAATGAATGAAT	G G G G G G G G G G G G G G G G G G G	I TTAAG K ATGAT D TAAGT S ATGTA V CCTTT F	K SAT' I TAT' I TAT' F AAGA R CGCA	N ITT L IAA K IGC A AGA D AGC A	AGA'D AACCTT TAAAK CTTC F	T TCC: P GGC: A AAATT I CACT T T TTCA S	G ITAT Y IAAA K IGCT A ITCT S AGAG E	3960 4020 4080 4140
TACT. Y Y GCAA A K GCAG A A AATT' N F GATCO D R AAAC' K L TATT'	ATT. Y AGTT F TTAM K GATC S TAGM D	V ATC L CAT L TTG V AAG G CTT L ATT Y ATG	TTTAC Y TAGCA A TAAAT N GAAGA TGGAT D	Q CGAA E AGAG CCCA P ACAA Q CGGA ACAG	T AT I TG W AG S GA D AA K AA K AT	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R R ACTT L V AAAA K GGA	AGGT G FAAT N FGGA I NAAT N AGTT V GTCA	AAGG K ACT T CCCT Y CAAA Q ACAA T CCCCC	K GAT D GTT V CAA Q GAA E TTT F CAC	ACACACA	G G G G G G G G G G G G G G G G G G G	TTAAGT ATGAT TAAGT S ATGTA V CCTTT F AGCTT	K SAT' I TAT' I TAT' F AGA R 'GCA A	N ITT L IAA K IGC A AGA D AGC A ACC P	AGA' D AACO T TAAA K CTTO F GGTA V	T TCCT P GGGCT A AATTT T TCACT S AATTG	G ITAT Y IAAA K IGCT A ITCT S AGAG E GAGT S	3960 4020 4080 4140 4200
TACT Y Y GCAA A K GCAG A A AATT N F GATC D R AAAC K L TATT Y F	ATT. Y AGTO S CTTTA K GATO S TAGA D TTTTA Y ACAA	V ATC L CAT L ITG V AAG G CTT L ATT Y ATT ATT Y ATT	TTTAC Y TAGCA A TAAAT N GAAGA R TGGAT D ATTTA	Q CGAA E AGAG CCA P ACAA Q ACAG G CGAA CGAA CTGAA CTTAT	T AT I TG W AG S GA D AA K AA M GA	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	L AAGA R FAGT S ACTT L CGTT V AAAA G G CAG	AGGT G FAAT N FGGA I AAAT N AGTT V GTCA S GAGC	TACCT P TACCY ACAA CGCC R TAT	K GAI D GTI V CAA Q GAA E TTT CAC TCA	AAATCAAAATCAAAATCAAAATCAAAATCAAAATCAAAATCAAAATCAAAAATCAAAAAA	G GV GD TL PH APP PA AP	TTAAGT ATGAT TAAGT S ATGTA V CCTTT F AGCTT L CTTAC	K SAT' I TAT' F AGGA R CGCA A TTA	N ITT L ITAA K IGC A AGA A ACC P TTC S	AGA' D AACO T TAAA K CTTO F CTTT C GGTA V CTCA	T TCC: P GGGC: A AAATT I CACT T TTCA S AATTCA L AGACA D TTCA	G ITAT Y IAAA K IGCT A ITCT S AGAG E GAGT S CAAT N	3960 4020 4080 4140 4200 4260 4320

AAACGTGGCA TGGGGGTTAT ACTTGATGTC GTCTATAATC ACACTGCAAA AACTTATCTC K R G M G V I L D V V Y N H T A K T Y L	4500
TTTGAGGATA TAGAACCTAA TTATTATCAC TTTATGAATG AAGATGGTTC ACCAAGAGAA F E D I E P N Y Y H F M N E D G S P R E	4560
AGTTTTGGAG GGGGACGTTT AGGAACCACT CATGCAATGA GTCGTCGTGT TTTGGTTGAT S F G G G R L G T T H A M S R R V L V D	4620
TCCATTAAAT ATCTTACAAG TGAATTTAAA GTTGATGGTT TCCGTTTTGA TATGATGGGA S I K Y L T S E F K V D G F R F D M M G	4680
GATCATGATG CGGCTGCGAT TGAATTAGCT TATAAAGAAG CTAAAGCTAT TAATCCTAAT D H D A A A I E L A Y K E A K A I N P N	4740
ATGATTATGA TTGGTGAGGG CTGGAGAACA TTCCAAGGCG ATCAAGGTCA GCCGGTTAAA M I M I G E G W R T F Q G D Q G Q P V K	4800
CCAGCTGACC AAGATTGGAT GAAGTCAACC GATACAGTTG GCGTCTTTTC AGATGATATT PADQ DWMKSTDTVGVFSDDI	4860
CGTAATAGCT TGAAATCTGG TTTTCCAAAT GAAGGTACTC CAGCTTTCAT CACAGGTGGC R N S L K S G F P N E G T P A F I T G G	4920
CCACAATCTT TACAAGGTAT TTTTAAAAAT ATCAAAGCAC AACCTGGGAA TTTTGAAGCA P Q S L Q G I F K N I K A Q P G N F E A	4980
GATTCGCCAG GAGATGTGGT GCAGTATATT GCTGCACATG ATAACCTTAC CTTGCATGAT D S P G D V V Q Y I A A H D N L T L H D	5040
GTGATTGCAA AATCAATT (SEQ ID NO:22) V I A K S I .	5058
FIG. 4a	
NLKAELSVED EQYTATVYGK SAHGSTPQEG VNGATYLALY LSQFDFEGPA	50
RAFLDVTANI IHEDFSGEKL GVAYEDDCMG PLSMNAGVFQ FDETNDDNTI	100
ALNFRYPQGT DAKTIQTKLE KLNGVEKVTL SDHEHTPHYV PMDDELVSTL	150
LAVYEKQTGL KGHEQVIGGG TFGRLLERGV AYGAMFPGDE NTMHQANEYM	200
PLENIFRSAA IYAEAIYELI K (SEQ ID NO:23)	221
FIG. 4b	
MTDLEKIIKA IKSDSQNQNY TENGIDPLFA APKTARINIV GQAPGLKTQE	50
ARLYWKDKSG DRLRQWLGVD EETFYHSGKF AVLPLDFYYP GKGKSGDLPP	
RKGFAEKWHP LILKEMPNVQ LTLLVGQYAQ KYYLGSSAHK NLTETVKAYK	
DYLPDYLPLV HPSPRNQIWL KKNPWFEKDL IVDLQKIVAD ILKD	194
(SEQ ID NO:24)	
FIG. 4c	

MRDNHLHTYF SYDCQTAFE	ED YINGFTGEFI	TTEHFDLSNP	YTGQDDVPDY	50
SAYCQKIDYL NQKYGNRFF	KK GIEIGYFKDR	ESDILDYLKN	KEFDLKLLSI	100
HHNGRYDYLQ EEALKVPTE	KG AFSRLL (SI	EQ ID NO:25)		126

FIG. 4d

MKRKDLFGDK	QTQYTIRKLS	VGVASVTTGV	CIFLHSPQVF	AEEVSVSPAT	50
TAIAESNINQ	VDNQQSTNLK	DDINSNSETV	VTPSDMPDTK	QLVSDETDTQ	100
KGVTEPDKAT	SLLEENKGPV	SDKNTLDLKV	APSTLQNTPD	KTSQAIGAPS	150
PTLKVANQAP	RIENGYFRLH	LKELPQGHPV	ESTGLWIWGD	VDQPSSNWPN	200
GAIPMTDAKK	DDYGYYVDFK	LSEKQRKQIS	FLINNKAGTN	LSGDHHIPLL	250
RPEMNQVWID	EKYGIHTYQP	LKEGYVRINY	LSSSSNYDHL	SAWLFKDVAT	300
PSTTWPDGSN	FVNQGLYGRY	IDVSLKTNAK	EIGFLILDES	KTGDAVKVQP	350
NDYVFRDLAN	HNQIFVKDKD	PKVYNNPYYI	DQVQLKDAQQ	IDLTSIQASF	400
TTLDGVDKTE	ILKELKVTDK	NQNAIQISDI	TLDTSKSLLI	IKGDFNPKQG	450
HFNISYNGNN	VMTRQSWEFK	DQLYAYSGNL	GAVLNQDGSK	VEASLWSPSA	500
DSVTMIIYDK	DNQNRVVATT	PLVKNNKGVW	QTILDTKLGI	KNYTGYYYLY	550
EIKRGKDKVK	ILDPYAKSLA	EWDSNTVNDD	IKTAKAAFVN	PSQLGPQNLS	600
FAKIANFKGR	QDAVIYEAHV	RDFTSDRSLD	GKLKNQFGTF	AAFSEKLDYL	650
QKLGVTHIQL	LPVLSYFYVN	EMDKSRSTAY	TSSDNNYNWG	YDPQSYFALS	700
GMYSEKPKDP	SARIAELKQL	IHDIHKRGMG	VILDVVYNHT	AKTYLFEDIE	750
PNYYHFMNED	GSPRESFGGG	RLGTTHAMSR	RVLVDSIKYL	TSEFKVDGFR	800
FDMMGDHDAA	AIELAYKEAK	AINPNMIMIG	EGWRTFQGDQ	GQPVKPADQD	850
WMKSTDTVGV	FSDDIRNSLK	SGFPNEGTPA	FITGGPQSLQ	GIFKNIKAQP	900
GNFEADSPGD	VVQYIAAHDN	LTLHDVIAKS	I (SEQ ID	NO:26)	931

FIG. 4e

1 Q 5	T TTGACAGAA L T E	G GTCAACTTC(G Q L R	S TTCTGATATC S D I	CCTGAGTTCC P E F F		60
> TACTGTACG' T V R	T GTTCACGCT V H A	A AAGTTGTTG <i>I</i> K V V E	AGGTACTCGC G T R	GAACGTATTC	AGATCTTTGA	120
		A AAGGTCAAGG		E R I Q		
G V V	1 5 K	K G Q G	I S E	M Y T V	RKI	180
TTCTGGTGGT S G G	I G V	G AGCGTACATI E R T F	CCCAATTCAC P I H	ACTCCTCGTG T P R V		240
CGAAGTTGTT E V V	CGTTATGGT R Y G	A AAGTACGTCG K V R R	TGCTAAACTT A K L	TACTACTTAC Y Y L R		300
AGGTAAAGCT	GCACGTATT	A AAGAAATCCG	TCGTTAATTT	TGATGATCAG	ATTTTAAAAA	360
		A GTAACTATGT				420
TCTGCCTGTG	G GGCAGTTTT	TTACTAGGTC	CCCTTAGTTC	AATGGATATA . H I Y		480
CCTAAGGAGI G L S	AATTGCTGGT Y N S	TCGATTCCGG	CAGGGGACAT C P V	ATTCATTGCA Y E N C	TGTAAATAGC T F L	540
GGTTTAGAGO P K S	TATTTTGCCC S N Q C	CAAATTTCTC L N R	TGATTAAGTT Q N L	TATCGTTCCT K D N R		600
TCTTGTAATT E Q L	GATGTGCGTA Q H A Y	AACTTCTAAA VEL	GTGATATTTA T I N	AATTCTCGTG L N E H	ATCTAAAACT D L V	660
TGAGAGATGG Q S I	AAATTAGATA S I L Y	GCTTGCAAAT S A F	GTATGCCTGA T H R	GAGAGTGCAC L S H V	TCGTACCTCG R V E	720
CGACCAGTTA R G T	TTTTTCGGAT	AGTTTTATTG T K N	ACTGCATTAT V A N	TTGAAAGTTT N S L K	GTCGAATAAT D F L	780
CTGTCGTTTT R D N	TATTTTTGT K N K T	AAATTCATGC F E H	AAAAAAAATA L F F	ATGTATCATT L T D N	GTCAATTGGT D I P	840
ATATTTCTGA I N R	TACTACTTTT I S S K	GTTTTTTGTT N K T		TTTGGTTGAA R Q N F	ATGATAATCC H Y D	900
CAAGTTTTAT W T K	TAATTGATAA N I S L	ATATTTGTTA Y K N	GTGTAATCAA T Y D	TATCATTAAC I D N V	TGTTAAACCT T L G	960
AAACATTCAG L C E		GCCAGTTTTA	GCGATGAGGT	ATAACGCTGC	_	1020
TGTTGTGATT	> TTTCTTTACA	AATTTTTATC	AAGCGTAAGT	ATTCATTGGT	TTCAAGAAAT	1080
		TTATTTTTTG				1140
		ACAGCCATGG				1200
		TGCGTTTGAC				1260

TGGAAGTAAT	ATTGCAAAGT	AATATATTTC	CTATTATAT	TTTATACGAT	ATTCGATATT	1320
CCCACCCGTT	GTCGCGTTTA	CGGAAATACG	CCATTGATAT	ACTCCACATT	AGCTAAAGAA	1380
CAGGGTGTTC	AAGGCTACCT	TGATGGAAAA	GGCTCTCTTA	GAGATATTTG	TAAATGGTAT	1440
GATATCTCAA	GTCGCTCTGT	TCTCCAAAAG	TGGATAAAAC	GGTATACTAG	TGGTGAAGAC	1500
TTGAAAGCCA	CTAGTAGAGG	ATATAGCCGT	ATGAAACAAG	GAAGGCAAGC	CACATTTGAA	1560
GAACGTGTAG	AGATTGTTAA	CTACACCATT	GCCCATGGGA	AAGACTATCA	AGCAGCTATT	1620
GAGAAGTTTG	GTGTTTCCTA	CCAACAAATT	TATTCTTGGG	TGCGTAAGCT	TGAGAAGAAT	1680
GGCTCACAAG	GTTTGGTTGA	TAGACGTGTG	AAAGGGTTGG	AGAGTAGGCC	TGATTTAACC	1740
GAGATTGAGC	AACTTTAACT	CAAGATTAAA	CAATTGGAGG	AACGTAATCG	TCTCTTAGAA	1800
ATCGAGGTTA	GTTTACTAAA	AAAGTTAGAA	GACATCAAAC	GAGGAAACAG	ACGGTAAGAC	1860
TAGGTAAGCA	TTTAGCGGAG	TTCCAAGTAA	TCAAGAATTA	TTACGATGAG	GAATCTAATG	1920
TGCCTATTCA	GGCCTTATGC	CAACTCTTGA	AGGGGTCTCG	TTCAGGCTAT	TACAAGTGGC	1980
TCAATCGTCA	AAAAACAGAT	TTTGAGACAA	AAAATACAAA	GCTAATGGCT	AAAATCAAGG	2040
AACTTCGTAG	ACTCTACAAT	GGTATCTTAG	GTTATCGCCG	TATGACAACA	TTTATTAATC	2100
GTCAACTTGG	GACAACTTAA	AACAAGAAAC	GGATTCGTTG	ATTGATGAAC	ATTCTGGGGA	2160
TTAGTTCAGT	CATTCGTCGT	GTTAGCCATG	CTTGTACAAA	AGCTGGTGAC	AGATTTTACG	2220
AAGAAAATAT	TCTTAATCGT	GAATTTACAG	CCACAGCTCA	TAACCAGAAA	TGGTGCACAG	2280
ATGTCACCTA	TCTTCAATAC	GGTCTGGGAG	CTAAAGCTTA	TCTCAGTGCG	ATTAAAGACC	2340
TGTATAACGG	TTCTATTATC	GCTTATGAGA	TTAGTCACAA	CAATGAAATC	CACTTGTTAT	2400
GAAGACCATT	AAAAAGGGGC	TAGAGCTCAA	TCCAGGAGCC	ACACCTATCA	TCCATAGCGA	2460
TTGAGGTAGT	CAATATACTT	CCAAAGAATA	CCGTTATATC	ATACAACAAG	CTGGTCTGAC	2520
CTTATCCATG	TCCCGGATTG	GCAAATGTAT	TGATAATGCA	CCAACTGAAA	GTTTCTTTGG	2580
	ACTGAGTCTT					2640
TGATGTGGCA	CGTTATATCG	AATTCTACAA	CACACAACGT	TATCAATCAA	AATTAAACAA	2700
CCTGACTCCT	CTAGAATTCA	GGAATCAGGT	TGCATAACTT	ATCTTTTATT	ATTTGACTGT	2760
CTACTTGACA	GGGAGCCGTT	CAGATTGCTT	AACCTTTCTA	AATTTGCTAA	AATAGCTACA	2820
AGAAAACGAG	CCATTTAATG	CTTATTTCTT	ATACTGTCTT	GCCTCACGCT	CTCCTCGACC	2880
AAAAATTGAG	CGTGAGGCTT	TTTGTTTCAT	TAAACGATGA	TATTTCCATA	TTCATCAGTT	2940
TGTTTTCCGA	GAGCCATCAA	AGCTTCGATA	AGGTCGATAA	TTCCAGGAAT	AAAGGTAATA	3000
СТАААААТАА	TATATAAAAA	AACCTGGCCT	ATTTTTCCTG	CGTAAAATTT	ATGCGCTCCA	3060
ATGCCGCCCA	AAAGAACGTT	AATAAAACAT	AAACTACTAT	GTTAGCATAA	GACTTTATTT	3120

TTACAACTG	A ATTTCATATA	AATGGATTA	G AGTAAGGGA	T AAAAGAAAT	AGCATAGCTC	3180
TTTTGAAAA	T AAAAAATTA	ATATAATAT	G GAAAAAATT	T TATTTCATA	A ACGTTTCATA	3240
AAAGGTATG	T AATCTAGTAT	'TTAGGCAAC	A CTATTTTGT	C ACTGGTGTC1	AGTAACTTAT	3300
AGATTGATA	A TTTTACTAGT	' AAACGTAAT1	CTTCGCTTT	A AGAGTTAAA1	GTCTATTTAT	3360
TGTAAGCTAA	ATTGGGAGGT	GAACTTATGT	T AAAATTAGA	T AGGTACTGTO	AAGTACGGGA	3420
TGATTATTGA	AACAGCCAGT	ATGCATCATA	A AAATCTGTA	T TGCTTAATAA	CTATTTCCTT	3480
AACCAGACAT	CAGTTCATTG	TTTATCATCO	CTACCCTAAC	G TCTAGTTTT	TCAATAGAGC	3540
ATTAGGTAGT	TTTTGATAAT	AAAACTATAT	AAACATGAG <i>I</i>	A ATTAGATTTC	GTATTGCATT	3600
CTTCATAATG	AGTTATTTGA	GATTTTCCTT	TGAATAAAT	GATACGAAAT	TCAGTAACTT	3660
CATATATAAA	CGGCTCTATC	ATTGAGATAG	TTTGTCAAA1	GAAGAAATTT	TTAATGGAAA	3720
TAGTTTTAAA	AACATTAGTT	GTAGGCGATG	: ТАААААТАТІ	AATCCAGTGG	ATGCAATAGT	3780
TGCGGAGTAA	AAATAGAGAG	GAGTAATTAG	GAAGTGATAA	AAAATGCTAT	AGCATATATT	3840
ACCAGAAAAA	AAAATAGAAC	ACTTATTATA	TTTGCTATTT	TAACAATTGT	TCTTTCTTGC	3900
TTGTATTCAT	GTTTAACAAT	AATGAAATCA M K S	AGTAATGAAA S N E I		TTTATATGAA L Y E	3960
AGTTCTAATT S S N S	CTTCAATATC S I S		AAAGATGGTA K D G K	AATATTTTAA Y F N	TATTAATCAA I N Q	4020
TTTAAGAATA F K N I	TTGAAAAAAT E K I	AAAAGAGGTT K E V	GAAGAAAAA E E K I		TGATGGATTA D G L	4080
GCAAAATTGA A K L K	AAGATCTTAA D L K	AGTAGTTAGT V V S	GGTGAGCAAA G E Q S		AGAAGATTTA E D L	4140
TCTGACGAAT S D E F	TTAAAAATGT K N V	TGTTTCACTA V S L	GAAGCTACAA E A T S	GTAATACTAA N T K	AAGAAATCTT R N L	4200
TTATTTAGTA L F S S	GTGGAGTATT G V F	TAGTTTTAAA S F K	GAAGGAAAA E G K N	ATATAGAAGA I E E	AAATGATAAG N D K	4260
AATTCAATTC N S I L	TTGTTCATGA V H E	AGAATTTGCT E F A	AAACAAAACA K Q N K		GGGTGATGAA G D E	4320
ATTGATCTTG I D L E	AATTACTAGA L L D	TACGGAAAAA T E K	AGTGGAAAA S G K I	TAAAAAGTCA K S H	TAAATTTAAA K F K	4380
ATTATAGGAA I I G I	TCTTTTCTGG F S G	TAAAAAACAG K K Q	GAAACATATA E T Y T	CAGGATTATC G L S	ATCTGATTTT S D F	4440
AGCGAAAATA S E N M	TGGTTTTTGT V F V	AGATTATTCA D Y S	ACTAGCCAAG T S Q E	AAATATTAAA I L N	TAAATCAGAG K S E	4500
AATAATAGAA N N R I	TTGCAAATAA A N K	AATTTTAATG I L M	TATTCTGGTA Y S G S		TACAGAGCTT T E L	4560
GCCTTAAACA	AATTGAAAGA	CTTTAAAATT	GATAAGTCAA	AGTATTCTAT	TAAGAAAGAT	4620

A L N K	L K D	F K I	D K S K	Y S I	K K D	
AATAAAGCAT N K A F	TCGAAGAGTC E E S		GTGAGTGGAA V S G I	TAAAACATAT K H I	AATTAAAATA I K I	4680
ATGACTTATT M T Y S	CGATTATGTT . I M L	AGGTGGAATA G G I	GTTGTTCTTT V V L S	CATTAATCTT L I L	GATTCTATGG I L W	4740
TTAAGAGAAA L R E R	GAATTTATGA A	AATAGGTATA I G I	TTTTTATCTA F L S I	TTGGAACAAC G T T	TAAGATACAA K I Q	4800
ATTATAAGGC I I R Q	AATTTATATT '		TTCATATCAA F I S I	TACCAAGTAT P S I	AATATCCTCC I S S	4860
TTATTTTAG L F L G	GGAATCTACT A		ATTGTAGAAG I V E G		CTCAGAGAAC S E N	4920
TCAATGATTT S M I F	TCGGTGGAAG 1 G G S		AAAAGCAGTT K S S F	TTATGTTAAA M L N	CATAACAACA I T T	4980
CTTGCAGAAA L A E S	GTTATTTAAT A		ATTATTGTTT I I V L	TATCAGTTGT S V V	AATGGCCTCT M A S	5040
TCATTAATAT S L I L	TATTTAAGAA A		ATATTATCAA I L S K	AAATAAGTTA I S .	GGAGCAAATA	5100
ATGGATATAT M D I L	TAGAAATAAA (E I K	GAATGTAAAT N V N	TACAGTTACG Y S Y A	CAAATTCTAA N S K	AGAAAAAGTT E K V	5160
TTGTCAGGAG L S G V	TAAATCAAAA A N Q K		GGAAAGTTTT G K F Y		AGGGAAGTCA G K S	5220
GGAACAGGAA G T G K	AATCCACACT S		CTTGCAGGAC L A G L	TTGATAAAGT D K V	TCAAACAGGA Q T G	5280
AAAATCTTGT K I L F	TTAAGAATGA A K N E		AAGAAAGGAT K K G Y	ATAGTAATCA S N H	CAGAAAAAAT R K N	5340
AATATATCTT N I S L	TGGTATTTCA A		TTAATAGATT L I D Y		GATTGAAAAT I E N	5400
ATTAGACTAG I R L V	TAAATAAATC A N K S		AGTATCTTGT S I L F	TCGAATTAGG E L G	TTTAGATAAA L D K	5460
AAACAAATAA K Q I K	AAAGAAATGT : R N V		TCTGGTGGTC S G G Q	AGCAACAAAG Q Q R	GGTAGCTATT V A I	5520
GCTAGGGCAC A R A L	TGGTATCAGA T		ATACTAGCTG I L A D	ATGAGCCTAC E P T	CGGTAACCTA G N L	5580
	CTGCTGGAGA A		SEQ ID NO:2	27)		5607

FIG. 5a

IQSLTEGQLR	SDIPEFRAGD	TVRVHAKVVE	GTRERIQIFE	GVVISRKGQG	50			
ISEMYTVRKI	SGGIGVERTF	PIHTPRVDKI	EVVRYGKVRR	AKLYYLRALQ	100			
GKAARIKEIR	R (SEQ ID	NO:28)			111			
		FIG. 5	db					
		LSINKTWDYH			50			
		KLSNNAVNKT			100			
		YAHQLQEQKD		GQNSSKPLFT	150			
CNEYVPCRNR	TSNYSLGGSC	YIH (SEQ	ID NO:29)		173			
		FIG. 5	C					
MKGGNETEKA	IVECONCCIC	TTEPOCEVEN	TNORWITHE					
		ITKKDGKYFN			50			
		EDLSDEFKNV			100			
		EFAKQNKLKL			150			
		SDFSENMVFV			200			
		FKIDKSKYSI			250			
		ILWLRERIYE			300			
		LKVIVEGFIN		LINKSSFMLN	350			
ITTLAESYLI	LISIIVLSVV	MASSLILFKK	PQEILSKIS		389			
(SEQ ID NO:	:30)	57. 0 5						
		FIG. 5	a					
MDILEIKNVN	YSYANSKEKV	LSGVNQKFEL	GKFYAIVGKS	GTGKSTLLSL	50			
		KKGYSNHRKN			100			
		KQIKRNVMKL			150			
ILADEPTGNL		(SEQ ID NO:			169			

FIG. 5e

CATATGACAA	TATTTTTCAA	AGTCTACATC	ACTTACTCGC	CTGTCGTGGA	AAATCTGGCA	60
ATACATTAAT	CGACCAATTA	GTTGCTGATG	GTTTACTTCA	TGCAGATAAT	CACTACCATT	120
TTTTCAATGG	GAAGTCTCTG	GCCACTTTCA	ATACTAACCA	ATTGATTCGC	GAAGTTGTCT	180
ATGTTGAAAT	ATCCTTAGAT	ACTATGTCTA	GTGGTGAACA	TGATTTAGTA	AAAGTTAACA	240
TTATCAGACC	CACTACCGAG	CATACTATCC	CCACGATGAT	GACAGCTAGC	CCCTATCATC	300
AAGGTATCAA	TGATCCTGCC	GCAGACCAAA	AAACATACCA	AATGGAGGGT	GCGCTAGCAG	360
TTAAACAGCC	TAAACACATA	CAAGTTGACA	CAAAACCATT	TAAAGAAGAA	GTAAAACATC	420
CTTCAAAATT	ACCCATCAGC	CCTGCAACTG	AAAGCTTCAC	ACACATTGAC	AGTTATAGTC	480
TCAATGACTA	TTTTCTTTCT	CGTGGTTTTG	CTAATATATA	CGTTTCAGGT	GTGGGTACTG	540
CTGGCTCTAC	GGGTTTCATG	ACCAGTGGGG	ATTACCAACA	AATACAAAGC	TTTAAAGCAG	600
TCATTGATTG	GTTAAATGGT	AAGGTTACTG	CATTCACAAG	TCATAAACGA	GATAAACAAG	660
TCAAGGCTGA	TTGGTCAAAC	GGCCTTGTAG	CAACCACAGG	TAAATCTTAT	CTCGGTACCA	720
TGTCAACTGG	TTTAGCAACA	ACTGGCGTTG	AGGGGCTGAA	AGTCATTATC	GCTGAAGCCG	780
CAATCTCCAC	ATGGTATGAT	TATTATCGAG	AAAATGGGCT	TGTGTGTAGT	CCAGGCGGCT	840
ACCCCGGTGA	AGATTTAGAC	GTTTTAACAG	AATTAACATA	CTCACGAAAC	CTCTTAGCTG	900
GTGATTACAT	CAAAAACAAC	GATTGCTATC	AAGCATTGTT	AAATGAACAA	TCAAAAGCAA	960
TTGACCGTCA	AAGTGGGGAT	TACAACCAAT	ACTGGCATGA	CCGTAATTAC	CTAACTCACG	1020
TCAATAATGT	CAAAAGTCGA	GTAGTTTACA	CTCATGGACT	ACAGGATTGG	AATGTTAAGC	1080
CAAGACATGT	CTACAAAGTT	TTCAATGCAT	TGCCTCAAAC	CATCAAAAA	CACCTTTTTT	1140
TACATCAAGG	TCAACATGTG	TATATGCATA	ATTGGCAGTC	GATTGATTTT	CGTGAAAGCA	1200
TGAATGCCTT	ACTAAGCCAA	GAACTACT1'G	GCATTGACAA	TCATTTCCAA	TTAGAAGAGG	1260
TCATTTGGCA	AGATAATACT	ACTGAGCAAA	CTTGGCAAGT	TTTAGATGCT	TTCGGAGGAA	1320
	GCAAATTGGT					1380
AAGAAGCCTT	TGATACTTAT	TGTAAAGACT	TCAATGTGTT	CAAAAATGAT	CTTTTCAAGG	1440
	AACCAATCAA					1500
	CAAACTCCAT					1560
CCCAAATCTT	AGACTATGGT	CCTAAAAAAC	GATTCAAAGA	TACACCAACC	ATCAAATTCT	1620
	TGATAATGGT					1680
	TTATCGTGTC					1740
	TGAGGCTATC					1800
	TCAATTGAGT					1860
	CATTCGAGAT					1920
	CCCAACTAAT					1980
	TCAACACTTT					2040
	CCATGATAAT					2100
	CGGAAGTTCG					2160
	TGCTCATTTA					2220
	CAATTTAATA					2280
	CGATACCCTA					2340
AACGCGAGGG	AGACTGATTA	ATGTCATCTT	ATTGGAATAA	CTATCCTGAA	CTTAAAAAAA	2400

ATATTGATGA	AACCAATCAA	CTAATTCAAG	AAAGAATACA	GGTCAGAAAT	AAAGATATTG	2460
AAGCGGCGCT	AAGCCAACTC	ACAGCTGCGG	GAGGAAAACA	GCTCAGACCA	GCATTCTTTT	2520
ACCTTTTTTC	TCAACTTGGT	AATAAGGAGA	ATCAAGATAC	TCAGCAACTA	AAGAAAATCG	2580
CTGCTTCTTT	AGAAATCCTT	CACGTTGCTA	CATTAATCCA	TGATGATGTC	ATTGATGACT	2640
CACCACTAAG	ACGTGGAAAT	ATGACCATTC	AAAGCAAGTT	TGGCAAAGAC	ATCGCAGTTT	2700
ATACTGGGGA	TTTACTTTTC	ACAGTCTTTT	TCGATCTTAT	TTTAGAATCT	ATGACTGATA	2760
CACCATTTAT	GAGGATTAAT	GCAAAATCTA	TGCGTAAAAT	TCTCATGGGA	GAATTGGACC	2820
AGATGCACCT	TCGTTACAAT	CAACAACAAG	GTATCCATCA	CTATTTACGT	GCGATTTCAG	2880
GTAAGACAGC	CGAACTCTTT	AAATTAGCTA	GCAAAGAAGG	AGCTTACTTT	GGTGGTGCAG	2940
AGAAGGAGGT	TGTTCGTCTA	GCAGGCCATA	TCGGCTTTAA	CATTGGTATG	ACATTCCAAA	3000
TTTTGGATGA	TATCCTGGAT	TATACTGCAG	ATAAAAAAAC	ATTTAATAAG	CCTGTCTTAG	3060
AGGATTTAAC	ACAAGGCGTT	TACAGCCTTC	CTCTACTTCT	TGCCATTGAA	GAAAATCCTG	3120
ATATTTTCAA	ACCTATTTTA	GATAAAAAA	CAGATATGGC	TACTGAAGAC	ATGGAAAAA	3180
TTGCTTATCT	CGTCGTTTCC	CATAGAGGTG	TTGACAAAGC	TCGCCATCTA	GCTCGTAAAT	3240
TTACTGAGAA	AGCTATTAGT	GACATAAATA	AGCTACCCCA	GAACTCTGCA	AAAAAACAGT	3300
TGCTACAATT	AACTAATTAC	CTTTTAAAAC	GCAAAATTTA	AATAATAAA	AAACATTCCA	3360
CAATGCTAGA	AAAGCAGTTA	GGGAATGTTT	TTTTATTATC	ATTTATTTAT	CGCACCTATC	3420
AATCATCATA	GATCACCATC	ATCAGCGGCT	TTCAGCTGAC	GGTAACGTTG	ACTACTTTGA	3480
GACAATTCTT	GAGGAGAACC	TTCCAACTCT	AATTGCCCAT	TTTCTATAAA	TAAGATACGA	3540
	CAATACCTTT		GTAATCCAAA	CTAAGGTCTT	ACCTTCCAAT	3600
	ATACCCTTAG	*		GATCAAGTCC		3660
			AGTAAGATTC	TAGCCAAAGC	AATTCTATGC	3720
	CTGAAAACCT		TCATCAACCA			3780
			CGTTCAAGAA		ATCTTCTTCA	3840
			TTTAGCAGGG		·	3900
GCTTGTTGTA	TCACTCCAAT	ATAGTTAGAA	ATGCAATCAC	CAACTATTGA	AACATCAGCA	3960
			TTCAAGTCGC	CACGAAGTAG	ACTAGCTAAG	4020
GTACTCTTGC	CAGAACCACT	CCGCCCTAAA	ATAGCAATTT	TTTCTCCTTC	TTTAATATCC	4080
	GATGCAAAAC		TGTGGCTTAT	ACTGGAAACT	TAAATTCTTG	4140
ACGGAAAAAT	CATATGGCTT	ATTAGGCAAT	T (SEQ ID	NO:32)		4171

FIG. 6a

PCT/CA99/00114 WO 99/42588

YDNIFQSLHH	LLACRGKSGN	TLIDQLVADG	LLHADNHYHF	FNGKSLATFN	50
TNQLIREVVY	VEISLDTMSS	GEHDLVKVNI	IRPTTEHTIP	TMMTASPYHQ	100
GINDPAADQK	TYQMEGALAV	KQPKHIQVDT	KPFKEEVKHP	SKLPISPATE	150
SFTHIDSYSL	NDYFLSRGFA	NIYVSGVGTA	GSTGFMTSGD	YQQIQSFKAV	200
IDWLNGKVTA	FTSHKRDKQV	KADWSNGLVA	TTGKSYLGTM	STGLATTGVE	250
GLKVIIAEAA	ISTWYDYYRE	NGLVCSPGGY	PGEDLDVLTE	LTYSRNLLAG	300
DYIKNNDCYQ	ALLNEQSKAI	DRQSGDYNQY	WHDRNYLTHV	NNVKSRVVYT	350
HGLQDWNVKP	RHVYKVFNAL	PQTIKKHLFL	HQGQHVYMHN	WQSIDFRESM	400
NALLSQELLG	IDNHFQLEEV	IWQDNTTEQT	WQVLDAFGGN	HQEQIGLGDS	450
KKLIDNHYDK	EAFDTYCKDF	NVFKNDLFKG	NNKTNQITIN	LPLKKNYLLN	500
GQCKLHLRVK	TSDKKAILSA	QILDYGPKKR	FKDTPTIKFL	NSLDNGKNFA	550
REALRELPFT	KDHYRVISKG	VLNLQNRTDL	LTIEAIEPEQ	WFDIEFSLQP	600
SIYQLSKGDN	LRIILYTTDF	EHTIRDNASY	SITVDLSQSY	LTIPTNQGN	649
(SEQ ID NO	:33)				
		FIG. 6	b		
MKLLTKERFD	DSQHFWYQIN	LLQESNFGAV	FDHDNKNIPQ	VVATIVDDLQ	50
GSGSSNHFWY	FGNTTDTSIL	MIAHLNRKFY	IQVNLKDFDF	ALNLIAINNW	100
KSLLQTQLEA	LNDTLAIFQ	(SEQ ID NO):34)		119
		FIG. 6	С		
MOGNETHING	T. (((0) T. D. T. (1)	-			
			KDIEAALSQL		50
			HVATLIHDDV		100
			MTDTPFMRIN		150
			KLASKEGAYF		200
			PVLEDLTQGV		250
			HRGVDKARHL	ARKFTEKAIS	300
DINKLPQNSA	KKQLLQLTNY	LLKRKI (SE	Q ID NO:35)		326

FIG. 6d

LPNKPYDFSV	KNLSFQYKPQ	EKWVLHHLDL	DIKEGEKIAI	LGRSGSGKST	50
LASLLRGDLK	ASQGKITLGG	ADVSIVGDCI	SNYIGVIQQA	PYLFNTTLLN	100
NIRIGNQDAS	EEDVWKVLER	VGLKEMVTDL	SDGLYTMVDE	AGLRFSGGER	150
HRIALARILL	KDVPIVILDE	PTVGLDPITE	QALLRVFMKE	LEGKTLVWIT	200
HHLKGIEHAD	RILFIENGQL	ELEGSPQELS	QSSQRYRQLK	AADDGDL	247
(SEQ ID NO:	:36)				

FIG. 6e

AATTCTATT	T GGAGGTTTTT	CTTGAATAAA	TGGTTAGTTA	AGGCAAGTTC	CTTAGTTGTT	60
TTAGGTGGT	TGGTTTTATC	TGCGGGTTCC	CGAGTTTTAG	G CGGATACTTA	TGTCCGTCCA	120
ATTGATAATO	G GTAGAATTAC	AACAGGTTTC	AATGGTTATO	CTGGACATTG	TGGGGTGGAT	180
TATGCTGTTC	CGACTGGAAC	GATTATTAGG	GCAGTGGCAG	ATGGTACTGT	GAAATTTGCA	240
GGAGCTGGAG	CCAACTTTTC	TTGGATGACA	GACTTAGCAG	GAAATTGTGT	CATGATTCAA	300
	GAATGCATAG					360
	AACAAGGAGA					420
CCTCACCTTC	ATTTTGAATT	TTTACCAGCT	AACCCTAATT	TTCAAAATGG	TTTCCATGGA	480
CGTATCAATC	CAACGTCACT	AATTGCTAAC	GTTGCGACCT	TTAGTGGAAA	AACGCAAGCA	540
TCAGCTCCAA	GCATTAAGCC	ATTACAATCA	GCTCCTGTAC	AGAATCAATC	TAGTAAATTA	600
AAAGTGTATC	GAGTAGATGA	ATTACAAAAG	GTTAATGGTG	TTTGGTTAGT	CAAAAATAAC	660
	CGACTGGGTT					720
	CTAATGGTAA					780
	CTAAAACTCT					840
	AGACACGCTT					900
	TTTACAAATA					960
	GTACAATTTC					1020
	GTCTCGGACT					1080
	ATTGACATCG					1140
	AGATACGACG					1200
	CAATAAATCA					1260
	GTCAATTGAT					1320
	TCCTGAGACA					1380
	AATAGAAACA					1440
	CAATCAAGTT					1500
	ACCAGAAGCA					1560
	GAAATCAAAA					1620
	GGTATCAACA					1680
	TAAACCAACT					1740
	TGAAACACCA					1800
	GGCAAGTGTT					1860
	AGCTCCAGCA					1920
	TGAAGTTAAG					1980
	AGCTTCAACA					2040
AACCTCATGT	TGCAGCTTAT	AAAGAAAAAG	TAGCGTCAAC	TTATGGAGTT	AATGAATTCA	2100
GTACATACCG	TGCAGGTGAT	CCAGGTGATC	ATGGTAAAGG	TTTAGCAGTC	GACTTTATTG	2160
TAGGTAAAAA	CCAAGCACTT	GGTAATGAAG	TTGCACAGTA	CTCTACACAA	AATATGGCAG	2220
CAAATAACAT	TTCATATGTT	ATCTGGCAAC	AAAAGTTTTA	CTCAAATACA	AATAGTATTT	2280
ATGGACCTGC	TAATACTTGG	AATGCAATGC	CAGATCGTGG	TGGCGTTACT	GCCAACCATT	2340
ATGACCATGT	TCACGTATCA	TTTAACAAAT	AAAAATATAA	AGGAAGCTAT	TTGGCTTCTT	2400

TTTTATATGC C	TTGAATAGA	CTTTCAAGGT	TCTTATCTAA	TTTTTATTAA	ATTGAGGAGA	2460
TTAAGCTATA A	GTCTGAAAC	TACTTTCACG	TTAACCGTGA	СТАААТСААА	ACGTTAAAAC	2520
TAAAATCTAA G	TCTGTAAAG	ATTATTGAAA	ACGCTTTAAA	AACAGATATA	ATAAGGTTTG	2580
TAGATATCTA A	AAAAATTAA	AGATAAGGAA	GTGAGAATAT	GCCACATCTA	AGTAAAGAAG	2640
CTTTTAAAAA G	CAAATAAAA	AATGGCATTA	TTGTGTCATG	TCAAGCTTTG	CCTGGGGAGC	2700
CTCTTTATAC TO	GAAAGTGGA	GGTGTTATGC	CTCTTTTAGC	TTTGGCAGCT	CAAGAAGCAG	2760
GAGCGGTTGG T	ATAAGAGCC	AATAGTGTCC	GCGACATTAA	GGAAATTCAA	GAAGTTACTA	2820
ATTTACCTAT C	ATCGGCATT	ATTAAACGTG	AATATCCTCC	ACAAGAACCA	TTTATCACTG	2880
CTACGATGAC A	GAGGTGGAT	CAATTAGCTA	GTTTAGATAT	TGCAGTAATA	GCCTTAGATT	2940
GTACACTTAG A	GAGCGTCAT	GATGGTTTGA	GTGTAGCTGA	GTTTATTCAA	AAGATAAAAG	3000
GGAAATATCC TO	GAACAGTTG	CTAATGGCTG	ATATAAGTAC	TTTTGAAGAA	GGTAAAAATG	3060
CTTTTGAAGC A	GGAGTTGAT	TTTGTGGGTA	CAACTCTATC	TGGATACACA	GATTACAGCC	3120
GCCAAGAAGA A	GGACCGGAT	ATAGAACTCC	TTAATAAGCT	TTGTCAAGCC	GGTATAGATG	3180
TGATTGCGGA A	GGTAAAATT	CATACTCCTA	AGCAAGCTAA	TGAAATTAAT	CATATAGGTG	3240
TTGCAGGAAT TO	GTAGTTGGT	GGTGCTATCA	CTAGACCAAA	AGAAATAGCG	GAGCGTTTCA	3300
TCTCAGGACT T	AGTTAAAAG	TGTTACTCAA	AAATCAAAAT	СААААТАААА	AAGGGGAATA	3360
GTTATGAGTA TO	CAAAAAAAG	TGTGATTGGT	TTTTGCCTCG	GAGCTGCAGC	ATTATCAATG	3420
TTTGCTTGTG TA	AGACAGTAG	TCAATCTGTT	ATGGCTGCCG	AGAAGGATAA	AGTCGAAATT	3480
(SEQ ID NO	:37)					

FIG. 7a

NSIWRFFLNK	WLVKASSLVV	LGGMVLSAGS	RVLADTYVRP	IDNGRITTGF	50
NGYPGHCGVD	YAVPTGTIIR	AVADGTVKFA	GAGANFSWMT	DLAGNCVMIQ	100
HADGMHSGYA	HMSRVVARTG	EKVKQGDIIG	YVGATGMATG	PHLHFEFLPA	150
NPNFQNGFHG	RINPTSLIAN	VATFSGKTQA	SAPSIKPLQS	APVQNQSSKL	200
KVYRVDELQK	VNGVWLVKNN	TLTPTGFDWN	DNGIPASEID	EVDANGNLTA	250
DQVLQKGGYF	IFNPKTLKTV	EKPIQGTAGL	TWAKTRFANG	SSVWLRVDNS	300
QELLYK (S	SEQ ID NO:38	3)			306

FIG. 7b

MKMNKKVLLT	STMAASLLSV	ASVQAQETDT	TWTARTVSEV	KADLVKQDNK	50
SSYTVKYGDT	LSVISEAMSI	DMNVLAKINN	IADINLIYPE	TTLTVTYDQK	100
SHTATSMKIE	TPATNAAGQT	TATVDLKTNQ	VSVADQKVSL	NTISEGMTPE	150
AATTIVSPMK	TYSSAPALKS	KEVLAQEQAV	SQAAANEQVS	TAPVKSITSE	200
VPAAKEEVKP	TQTSVSQSTT	VSPASVAAET	PAPVAKVAPV	RTVAAPRVAS	250
VKVVTPKVET	GASPEHVSAP	AVPVTTTSTA	TDSKLQATEV	KSVPVAQKAP	300
TATPVAQPAS	TTNAVAAHPE	NAGLQPHVAA	YKEKVASTYG	VNEFSTYRAG	350
DPGDHGKGLA	VDFIVGKNQA	LGNEVAQYST	QNMAANNISY	VIWQQKFYSN	400
TNSIYGPANT	WNAMPDRGGV	TANHYDHVHV	SFNK (SEQ	ID NO:39)	434
		FIG. 7	С		
MPHLSKEAFK	KQIKNGIIVS	CQALPGEPLY	TESGGVMPLL	ALAAQEAGAV	50
GIRANSVRDI	KEIQEVTNLP	IIGIIKREYP	PQEPFITATM	TEVDQLASLD	100
IAVIALDCTL	RERHDGLSVA	EFIQKIKGKY	PEQLLMADIS	TFEEGKNAFE	150
AGVDFVGTTL	SGYTDYXRQE	EGPDIELLNK	LCQAGIDVIA	EGKIHTPKQA	200
NEINHIGVAG	IVVGGAITRP	KEIAERFISG	LS (SEQ ID	NO:40)	232
		FIG. 7	d		
MSIKKSVIGF	CLGAAALSMF	ACVDSSQSVM	AAEKDKVEI		39

FIG. 7e

(SEQ ID NO:41)

ATGAAAATGA		ACTATTGACA	TCGACAATGG	CAGCTTCGCT	50
ATTATCAGTC	GCAAGTGTTC	AAGCACAAGA	AACAGATACG	ACGTGGACAG	100
CACGTACTGT	TTCAGAGGTA	AAGGCTGATT	TGGTAAAGCA	AGACAATAAA	150
TCATCATATA	CTGTGAAATA	TGGTGATACA	CTAAGCGTTA	TTTCAGAAGC	200
AATGTCAATT	GATATGAATG	TCTTAGCAAA	AATTAATAAC	ATTGCAGATA	250
TCAATCTTAT	TTATCCTGAG	ACAACACTGA	CAGTAACTTA	CGATCAGAAG	300
AGTCATACTG	CCACTTCAAT	GAAAATAGAA	ACACCAGCAA	CAAATGCTGC	350
TGGTCAAACA	ACAGCTACTG	TGGATTTGAA	AACCAATCAA	GTTTCTGTTG	400
CAGACCAAAA	AGTTTCTCTC	AATACAATTT	CGGAAGGTAT	GACACCAGAA	450
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TTTGAAATCA	AAAGAAGTAT	TAGCACAAGA	GCAAGCTGTT	AGTCAAGCAG	550
CAGCTAATGA	ACAGGTATCA	ACAGCTCCTG	TGAAGTCGAT	TACTTCAGAA	600
GTTCCAGCAG	CTAAAGAGGA	AGTTAAACCA	ACTCAGACGT	CAGTCAGTCA	650
GTCAACAACA	GTATCACCAG	CTTCTGTTGC	CGCTGAAACA	CCAGCTCCAG	700
TAGCTAAAGT	AGCACCGGTA	AGAACTGTAG	CAGCCCCTAG	AGTGGCAAGT	750
GTTAAAGTAG	TCACTCCTAA	AGTAGAAACT	GGTGCATCAC	CAGAGCATGT	800
ATCAGCTCCA	GCAGTTCCTG	TGACTACGAC	TTCAACAGCT	ACAGACAGTA	850
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ACAGCAACAC	CGGTAGCACA	ACCAGCTTCA	ACAACAAATG	CAGTAGCTGC	950
ACATCCTGAA	AATGCAGGGC	TCCAACCTCA	TGTTGCAGCT	TATAAAGAAA	1000
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GATCCAGGTG	ATCATGGTAA	AGGTTTAGCA	GTCGACTTTA	TTGTAGGTAA	1100
AAACCAAGCA	CTTGGTAATG	AAGTTGCACA	GTACTCTACA	CAAAATATGG	1150
CAGCAAATAA	CATTTCATAT	GTTATCTGGC	AACAAAAGTT	TTACTCAAAT	1200
ACAAATAGTA	TTTATGGACC	TGCTAATACT	TGGAATGCAA	TGCCAGATCG	1250
TGGTGGCGTT	ACTGCCAACC	ATTATGACCA	TGTTCACGTA	TCATTTAACA	1300
AATAA					1305

(SEQ ID NO:42)

FIG. 8

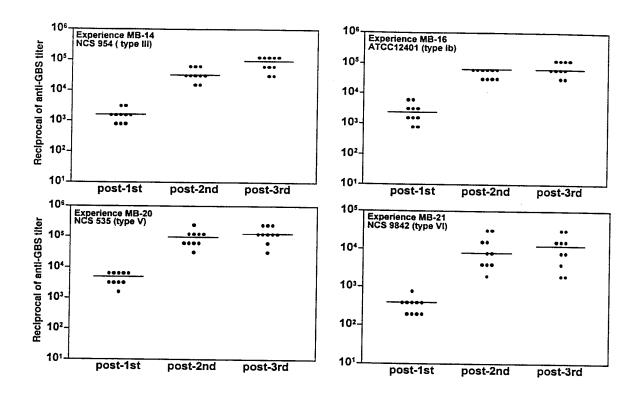
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GCAAAAATTA		AGATATCAAT	CTTATTTATC	CTGAGACAAC	200
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AAACTGGTGC	ATCACCAGAG	CATGTATCAG	CTCCAGCAGT	TCCTGTGACT	700
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CTGGCAACAA	AAGTTTTACT	·	AATAACATTT	CATATGTTAT	1100
ATACTTGGAA	TGCAATGCCA	CAAATACAAA	TAGTATTTAT	GGACCTGCTA	1150
GACCATGTTC	ACGTATCATT	GATCGTGGTG	GCGTTACTGC	CAACCATTAT	1200
GACCAIGIIC	ACGIATCATT	TAACAAATAA	(SEQ ID	NO:43)	1230

FIG. 9

QETDTTWTAR	TVSEVKADLV	KQDNKSSYTV	KYGDTLSVIS	EAMSIDMNVL	50
AKINNIADIN	LIYPETTLTV	TYDQKSHTAT	SMKIETPATN	AAGQTTATVD	100
LKTNQVSVAD	QKVSLNTISE	GMTPEAATTI	VSPMKTYSSA	PALKSKEVLA	150
QEQAVSQAAA	NEQVSTAPVK	SITSEVPAAK	EEVKPTQTSV	SQSTTVSPAS	200
	KVAPVRTVAA				250
	QATEVKSVPV				300
	ASTYGVNEFS				350
	NNISYVIWQQ			DRGGVTANHY	400
DHVHVSFNK	(SEQ ID NO:				
DITTITUDE	(SEQ ID NO:	44)			409

FIG. 9a

Fig. 10



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SEQUENCE LISTING

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            BRODEUR, Bernard R.
            HAMEL, Josée
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            BOYER, Martine
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15

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					ctt Leu 85											287
					ttt Phe											335
					tta Leu					-	_	_		-	_	383
					atg Met								_	_		431
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					gca Ala	Thr		Thr					Phe			590
					tta Leu											638
					gaa Glu 195											686
					gta Val											734

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	aag Lys 440															1569
	ttg Leu															1617
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gag Glu 530 tta Leu gac Asp att Ile	caa Gln att Ile ttt Phe	cta Leu atc Ile gta Val cag Gln 580	aag Lys ctt Leu ctg Leu 565 act Thr	agt Ser gtg Val 550 aat Asn agt Ser	gtt Val 535 ggg Gly gga Gly tgg Trp	ttt Phe gtt Val ctt Leu tgt Cys	ggg Gly atc Ile tta Leu atc Ile 585	caa Gln gct Ala cgt Arg 570 aac Asn	tta Leu gtc Val 555 aca Thr act	tct Ser 540 tta Leu gat Asp ttt Phe	cca Pro ccg Pro aaa Lys aat Asn	atg Met aca Thr agc Ser aac Asn 590	aat Asn acc Thr aaa Lys 575 ttg Leu	ctt Leu gga Gly 560 agg Arg tca Ser	ttc Phe 545 tat Tyr tat Tyr gga Gly	1918 1966 2014

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	aaa Lys 675															2350
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	ata Ile															2446
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9 / 63

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Val	Thr 370	His	Thr	Phe	Leu	Gly 375		Arg	Ile	Lys	Ala 380	Tyr	Gly	Lys	Gly
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Glu	Ser	Ile	His	Ser 405	Val	Asp	Lys	Ser	Gly 410	Val	Thr	Ala	Lys	His 415	Gly
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			740					745					750		
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345

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Pro Arg Leu Ala Val Asp Val Ser Ser Leu Pro Met His Ala Gly Asn
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Ala Thr Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His
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Val Lys Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser
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Gly Tyr Ile Phe Asp Pro Arg Asp Val Leu Ala Lys Glu Thr Phe Val
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Arg Thr Ile Asn Lys Ser Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala
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Lys Pro Lys Glu Lys Gln Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln
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Pro Ser Glu Ala Ser Lys Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp
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Ser Leu Pro Asp Tyr Gly Leu Asp Arg Ala Thr Leu Glu Asp His Ile
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Asn Gln Leu Ala Gln Lys Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe
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Thr Leu Ser Asp His Glu His Thr Pro His Tyr Val Pro Met Asp Asp
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(57) Abstract

Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.

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A. CLASSIF IPC 6	FICATION OF SUBJECT MATTER C12N15/31 C07K14/315 A61K39/0	09 C12N1/21	
According to	International Patent Classification (IPC) or to both national classification	tion and IPC	
· · · · · · · · · · · · · · · · · · ·	SEARCHED		
Minimum do IPC 6	cumentation searched (classification system followed by classificatio CO7K C12N A61K	n symbols)	
Documentat	ion searched other than minimum documentation to the extent that su	ich documents are included in the fields sea	arched
Electronic da	ata base consulted during the international search (name of data bas	e and, where practical, search terms used)	
C. DOCUME	ENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the rele	vant passages	Relevant to claim No.
A	MICHEL J L ET AL: "Cloned alpha C-protein antigens of group B Strelicit protective immunity" INFECTION AND IMMUNITY., vol. 59, no. 6, June 1991 (1991-62023-2028, XP002107260 AMERICAN SOCIETY FOR MICROBIOLOGY WASHINGTON., US ISSN: 0019-9567 the whole document	reptococci 96), pages	1-48
X Furt	her documents are listed in the continuation of box C.	Patent family members are listed	in annex.
° Special of 'A' docume consider 'E' earlier of filing of 'L' docume which citatio 'O' docume other 'P' docume later to 'P' docume of 'P' docume of	ernational filing date the application but eory underlying the claimed invention t be considered to comment is taken alone claimed invention ventive step when the ore other such doou- us to a person skilled family		
Date of the	actual completion of the international search	Date of mailing of the international sea	arch report
1	.5 December 1999	24 01 2000)
Name and	mailing address of the ISA European Patent Office, P.B. 5818 Patentiaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Lejeune, R	

PCT/CA 99/00114

		PC1/CA 99/00114
C.(Continu	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	To de contra de Pala
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	LACHENAUER C S ET AL: "Cloning and expression in Escherichia coli of a protective surface protein from type V group B Streptococci" ADVANCES IN EXPERIMENTAL MEDICINE AND BIOLOGY, vol. 418, 9 December 1997 (1997-12-09), pages 615-618, XP002107261 SPRING ST., NY, US ISSN: 0065-2598 the whole document	1-48
P,X	DATABASE EMBL [Online] Accession number AF062533, 11 February 1999 (1999-02-11) SPELLERBERG B ET AL: "Streptococcus agalactiae Lmb (lmb) gene, complete cds; and unknown gene." XP002125180 98.9% identity between base 1-2514 of SEQ ID NO 13 and base 988-3501 of AF062533 Translation product (AC: Q9ZHG9) has 98.5% identity in 793 AA overlap with SEQ ID NO 15 and 98.5% identity in 715 AA overlap with SEQ ID 16 & SPELLERBERG B ET AL: "Lmb, a protein with similarities to the LraI adhesin family, mediates attachment of Streptococcus agalactiae to human laminin" INFECTION AND IMMUNITY., vol. 67, no. 2, February 1999 (1999-02), pages 871-878, AMERICAN SOCIETY FOR MICROBIOLOGY. WASHINGTON., US ISSN: 0019-9567	1-10, 16-23,26
X	DATABASE EMBL [Online] Accession Number L23843, 4 January 1994 (1994-01-04) MACRINA F L ET AL: "ISN IS199 from Streptococcus mutans IS3 (Brathall serotype C) DNA fragment" XP002125181 79.6% identity between base 5212-4314 of SEQ ID NO 13 and base 312-1220 of L23843 Translation has 83.4% identity in 283 AA overlap with SEQ ID NO 21 -/	1,3-7,10

Interi nal Application No PCT/CA 99/00114

Category °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE EMBL [Online] Accession Number AF026542, 15 October 1997 (1997-10-15) HYNES W L ET AL: "Streptococcus pyogenes FF22 lantibiotic (scn) gene cluster region containing: scnK, scnR, streptococcin A-FF22 precursor (scnA), scnA1, scnM, scnT, scnF, scnE, scnG genes, complete cds, and tnpA gene, partial cds." XP002125182 88.2% identity between base 2607-2953 of SEQ ID NO 13 and base 10435-10777 of AF026542 Translation product (AC: 031057) has 95.8% identity in 71 AA overlap with SEQ ID NO 17	1-10, 16-23,26
P,X	DATABASE GENESEQ [Online] Accession Number V52136, 23 October 1998 (1998-10-23) BARASH S C ET AL: "Streptococcus pneumoniae genome fragment SEQ ID NO:3" XP002125183 68.5% identity between base 2539-3319 of SEQ ID NO 37 and base 18492-19271 of V52136 Translation has 74.5% identity in 231 AA overlap with SEQ ID NO 40 & WO 98 18931 A (DOUGHERTY BRIAN A ;HUMAN GENOME SCIENCES INC (US); ROSEN CRAIG A) 7 May 1998 (1998-05-07)	1,3-7,10

ational application No. PCT/CA 99/00114

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Box I	Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)
This Inte	ernational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:
1. X	Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
	Although claims 37-46 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2.	Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3.	Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box II	Observations where unity of invention is lacking (Continuation of item 2 of first sheet)
This Inte	ernational Searching Authority found multiple inventions in this international application, as follows:
	see additional sheet
	As a result of the prior review under R. 40.2(e) PCT, no additional fees are to be refunded.
1.	As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.	As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
з. 🛛	As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
	11-14,16,24,25,27,28,30,31 (completely), 1-10,15,17-23,26,29,32-48 (all partially) i.e. (group of) inventions 1, 3 and 7
4.	No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remari	con Protest X The additional search fees were accompanied by the applicant's protest.
	No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

An isolated polynucleotide encoding a polypeptide having a sequence selected from the group consisting of SEQ ID 2, SEQ ID NO 3, SEQ ID NO 4, SEQ ID NO 5, SEQ ID NO 6 i.e. the open reading frames of clone 1 (SEQ ID NO 1). Also a vector comprising the polynucleotide, a host cell transformed therewith, an isolated polypeptide encoded by the polynucleotide, a vaccine composition comprising said polypeptide and a polynucleotide having a sequence SEQ ID NO 1.

2. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 2 (SEQ ID 7) with sequences SEQ ID NO 8-12.

3. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 3 (SEQ ID 13) with sequences SEQ ID NO 14-21.

4. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 4 (SEQ ID 22) with sequences SEQ ID NO 23-26.

5. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 5 (SEQ ID 27) with sequences SEQ ID NO 28-31.

6. Claims: 1-10,15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 6 (SEQ ID 32) with sequences SEQ ID NO 33-36.

7. Claims: 11-14,16,24,25,27,28,30,31 (all completely), 1-10, 15,17-23,26,29,32-48 (all partially)

Same as invention 1, but directed at polypeptides of clone 7 (SEQ ID 37) with sequences SEQ ID NO 38-41.

information on patent family members

Interr hal Application No PCT/CA 99/00114

Patent document cited in search report	Publication date		Patent family member(s)	Publication date
WO 9818931 A	07-05-1998	AU AU EP EP WO	5194598 A 6909098 A 0942983 A 0941335 A 9818930 A	22-05-1998 22-05-1998 22-09-1999 15-09-1999 07-05-1998